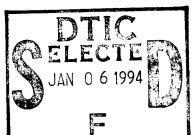


# FINAL REPORT FOR SOURCE TEST MEASUREMENT OF NITROGEN OXIDES, SULFUR DIOXIDE, CARBON MONOXIDE, VOC, AND PM10 EMISSIONS ON GAS TURBINE #2 AT ONIZUKA AIR FORCE BASE SUNNYVALE, CALIFORNIA

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2402 E Drive
Brooks Air Force Base, TX 78235-5114

**FEBRUARY 1994** 

Final Contractor Report for Period December 1993 - January 1994

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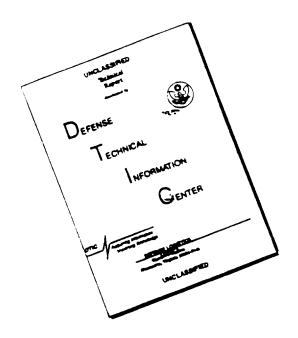
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#### **FOREWORD**

This report was prepared by Pacific Environmental Services, Inc. (PES) using due and ordinary care and in accordance with the scope of work identified in the Professional Services Agreement, Purchase Order, or other written/verbal request.

Any measured contaminants and their concentrations detected or identified in this study are for the sample(s) obtained or for the observations made at the time(s) and location(s) referenced in the report and may not represent other times, locations, or contaminants and their concentrations.

Except for the tests and observations conducted by PES, no attempt was made to check for compliance of present or past owners or operators of the equipment, plant, or site with federal, state, or local laws and regulations.

The information provided in this report, including any drawings and specifications, was prepared solely for the use of the identified client and any use by any other party shall be at their own risk.

The project work was conducted by Steven M. Hernandez, Robert T. Nguyen, and S. Hugh Brown under the direction of S. Hugh Brown.

Approved:

S. Hugh Brown, Director Air Quality Testing

5 Hugh Brown

A-1

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#### INTRODUCTION

Onizuka Air Force Base currently has twelve gas turbine generators which supply electric power for the entire facility. Heat recovery boilers are connected to each generator. Current plans are for the power plant to curtail its operation. Under current Bay Area Air Quality Management District (BAAQMD) rules and regulations, emissions credits can be banked and sold. Source testing is required to quantify the baseline emissions for each constituent potentially reduced.

Based on information obtained from the BAAQMD, source testing was conducted for nitrogen oxides (NO<sub>x</sub>), sulfur dioxide (SO<sub>2</sub>), carbon monoxide (CO), volatile organic compounds (VOC), and particulate matter less than 10 microns (PM10).

Pacific Environmental Services (PES), a participant in CARB's Independent Contractor Program, was hired by the Air Force to perform the required source testing and data reduction. PES qualifies as an independent testing laboratory (no conflict of interest). The source testing was conducted by S. Hugh Brown, Steven Hernandez, and Robert Nguyen of PES on December 3, 1993 and January 12, 1994.

### EQUIPMENT AND PROCESS DESCRIPTION

All of the gas turbine generators were identical 750 kVA units that were manufactured by Solar and each consumed a maximum of 130 therms/hour of natural gas. Gas turbine #2 was equipped with sampling ports and a work platform and was used as a demonstration unit for all twelve generators. The turbine exhaust gases were routed under the turbine platform and over to the heat recovery boiler adjacent to the turbine on the same platform. The gases entered the boiler through a diverter/bypass damper at the base of the exhaust stack. When the damper was in the bypass position, the gases were routed straight up the stack to atmosphere. When the damper was in the closed position, the gases were routed horizontally through the boiler exiting the top and rejoining the stack just above the bypass damper. The location of the turbines on the base is shown in Figure 2.1. The gas turbine and heat recovery boiler are depicted in Figure 2.2. A copy of the Permit to Operate issued by the BAAQMD is located in Appendix A.

The source testing was conducted on gas turbine #2 at a constant load of about 570 KW with the heat recovery boiler damper set at 75% closed (25% bypass).

Figure 2.1
Plant Layout And Equipment Location

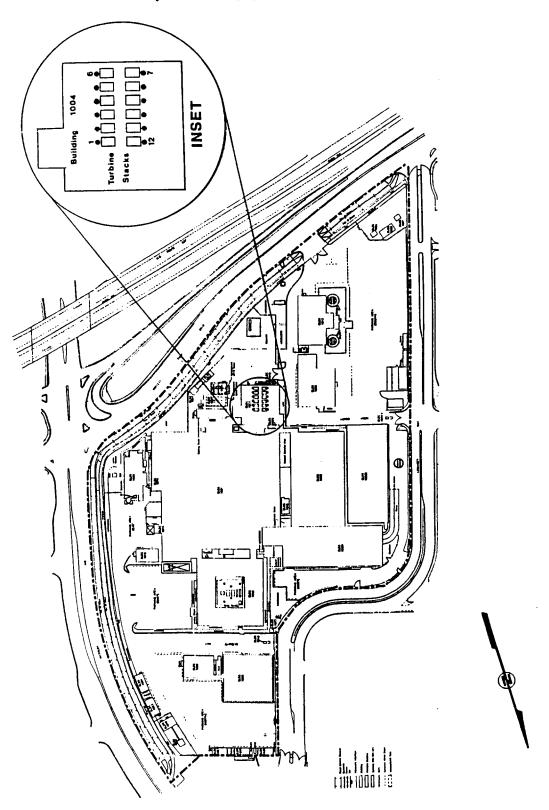
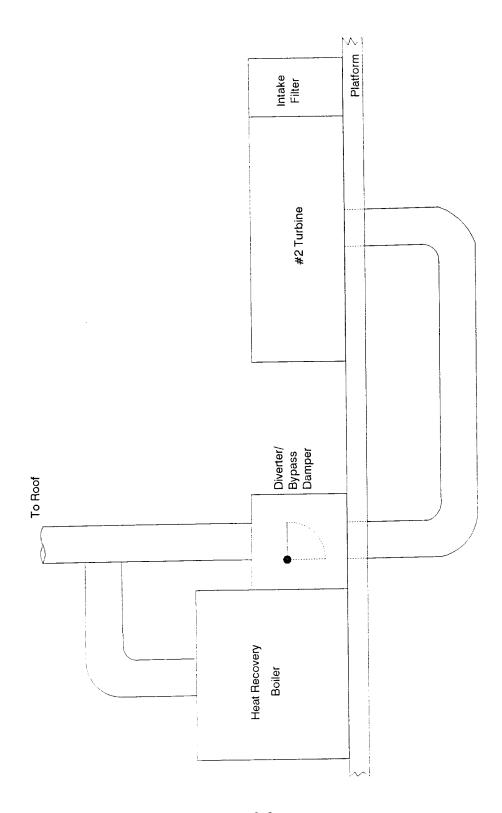


Figure 2.2 Schematic - Turbine/Heat Recovery Boiler Number 2



#### **TESTING METHODOLOGY**

The approximate sampling port locations are shown on Figure 3.1. The number of traverse points required (4 on each of two diameters, 90 degrees apart) and their locations were derived from CARB Method 1. The sampling was conducted on the exhaust of turbine #2 at roof level during block load conditions. Triplicate tests were conducted for each constituent determined.

#### PM10

The PM10 emissions of the exhaust were determined by EPA Method 201A. The sampling train is shown in Figure 3.2 and consisted of a stainless steel nozzle, a 10-micron cut-off cyclone and an in-stack stainless steel filter holder manufactured by Andersen, a 48-inch glass probe, a 10-foot Teflon hose from the probe to the first impinger, two Greenburg-Smith impingers each charged with 100 mls of distilled water, an empty impinger, an impinger filled with silica gel, a 30-foot umbilical line, a vacuum pump, a dry gas meter and a calibrated orifice connected to an oil inclined manometer. Glass fiber filters were used in the filter holders. Each sample was collected isokinetically at a fixed sampling rate and the number of minutes sampled at each traverse point was determined by the ratio of the point velocity to the average velocity. The velocity and temperature were measured at each traverse point as it was sampled. Field data and calculation sheets are included in Appendix B.

The volume of the impinger solution and the weight of the silica gel were recorded before and after the tests in order to obtain the moisture content of the stack gas stream. All sample volumes and weights were recorded immediately on sample recovery sheets (Appendix B) during charging and sample recovery. Leak checks were performed before and after each test. The post-test leak check was performed after removing the cut-off cyclone so as not to disturb the particle catch.

After the test, the contents of the nozzle and cut-off cyclone (PM10+) were recovered by rinsing three times with acetone. The washings were placed in a 125-milliliter polyethylene container. The filter was placed in a plastic cassette. The contents of the cyclone exit and filter holder front (PM10) were also recovered by rinsing three times with acetone. The washings were placed in a separate 125-milliliter polyethylene container. The contents of each impinger set was placed in a 1000-milliliter polyethylene container. The sampling train was then rinsed from the 3rd impinger to the nozzle with the distilled water and the rinse was added to the sample. The impinger solution was saved but not analyzed. Disposable vinyl gloves were worn during sample retrieval to help prevent contamination.

Figure 3.1 Sampling Location - Turbine Number 2

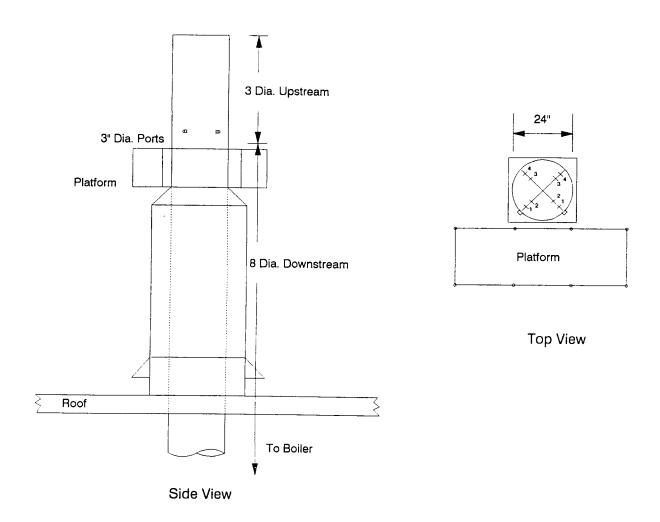
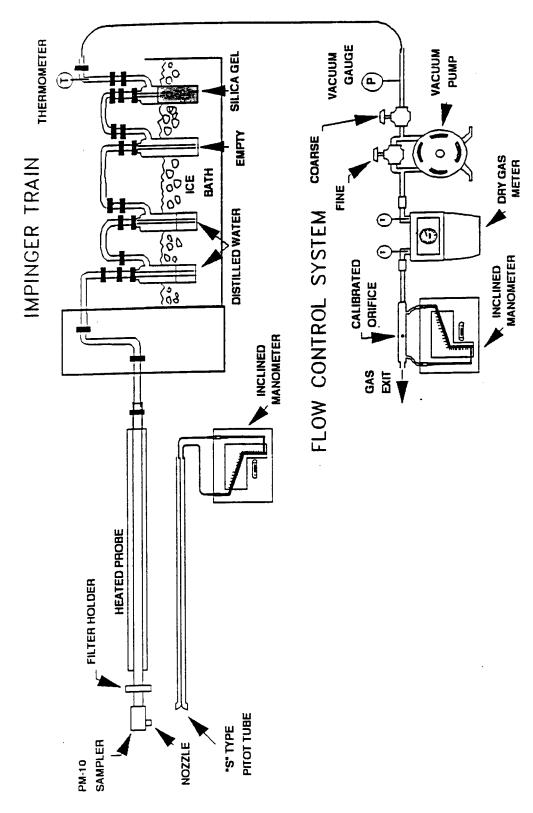


Figure 3.2
PM10 Sampling Train - EPA Method 201A



The particulate fractions were evaporated to dryness at 100 degrees Centigrade and desiccated to a constant weight along with the sample filter.

#### CONTINUOUS MONITORING

The turbine/boiler exhaust was monitored for VOC, CO, CO<sub>2</sub>, O<sub>2</sub> and NO<sub>x</sub> by CARB Method 100. Each test run was 60 minutes long. Rosemount Analytical Model 880 Infrared Analyzers were used to determine the CO and CO<sub>2</sub> concentrations. A Rosemount Analytical Model 755R Paramagnetic Analyzer was used to determine the oxygen concentration, and a Thermo Electron Model 10 Chemiluminescent Analyzer was used to determine the NO<sub>x</sub> concentration. The output of the analyzers was linearized by the manufacturers. Table 3.1 lists the instrument specifications. Table 3.2 lists the calibration gases that were used specific to this job.

The continuous monitoring train for the above gases is shown in Figure 3.3 and consisted of a 3/8-inch stainless steel sampling probe, a 3/8-inch heated Teflon sampling line, a sample refrigeration/pump system, a glass fiber filter in a 47-millimeter stainless steel holder, and a sample distribution manifold. The distribution manifold was equipped with a series of 3-way valves with flow meters (rotometer style). One flow meter acted as a bypass, and the others were connected to the individual analyzers. The output of the analyzers was logged by a Yokogawa Model HR2400 multi-channel recorder and a Rustrak Ranger II data logger.

The  $NO_x$  analyzer was operated on a range of 0-50 ppmv with span gases at 10.4, 23.5, and 44.2 ppmv. The CO analyzer was operated on a range of 0-100 ppmv with span gases at 50 and 75 ppmv. The  $O_2$  analyzer was operated on a range of 0-25 per cent with span gases at 11.0 and 19.0 per cent. The  $CO_2$  analyzer was operated on a range of 0-5 per cent with span gases at about 2.0 and 4.0 per cent. Prior to the source tests, the suction side of the monitoring system was leak-checked at a full vacuum (greater than 20 inches mercury).

A bias check was made on each analyzer by comparing the response between the span gas introduced at the sample line tip and the span gas introduced directly to the analyzer to ensure a differential of less than 5 per cent. The analyzers were spanned before and after each test run with NIST traceable calibration gases from Scott Specialty, and with zero grade nitrogen.

The total hydrocarbons were determined on January 12, 1994 by using a JUM Model VE-7 total hydrocarbon analyzer that utilized a heated Teflon sample line and a flame ionization detector mounted in a heated oven. The sample line was maintained at 250 degrees Fahrenheit, and the oven was maintained at 180 degrees Centigrade. Since the gas stream was expected to have a low concentration of

# Table 3.1 Continuous Monitoring Specifications

NO <sub>x</sub> Chemiluminescent Analyzer - T	hermo-Electron Model 10A
Response Time	1.5 sec - NO, 1.7 sec - NO <sub>X</sub>
Zero Drift	$\pm$ 0.5% after warm up (30 min)
Span Drift	+ 1% of full scale
Linearity	± 1% of full scale
Accuracy	Derived from the calibration $NO/NO_x$ $\pm$ 1% gas was used.
Output	NO 0-5.0 Vdc (scaled 0-50 ppm) NO <sub>x</sub> 0-0.5 Vdc (scaled 0-50 ppm)
O2 Paramagnetic Analyzer - Beckma	n Model 755R
Response Time	2 Sec
Zero Drift	± 1% of full scale
Span Drift	+ 1% of full scale
Linearity	± 1% of full scale
Accuracy	Derived from the calibration O2 <u>+</u> 1% gas was used.
Output	0-1.0 Vdc (scaled 0-25%)
CO/CO2 Infrared Analyzers - Beckn	nan Model 880
Response Time	2 sec
Zero Drift	+ 1% of full scale
Span Drift	+ 1% of full scale
Linearity	± 1% of full scale
Accuracy	Derived from the calibration CO ± 1% gas was used.
Output	0-1.0 Vdc (scaled 0-100 ppm CO) 0-1.0 Vdc (scaled 0-20 % CO <sub>2</sub> )

Table 3.1 - Continued

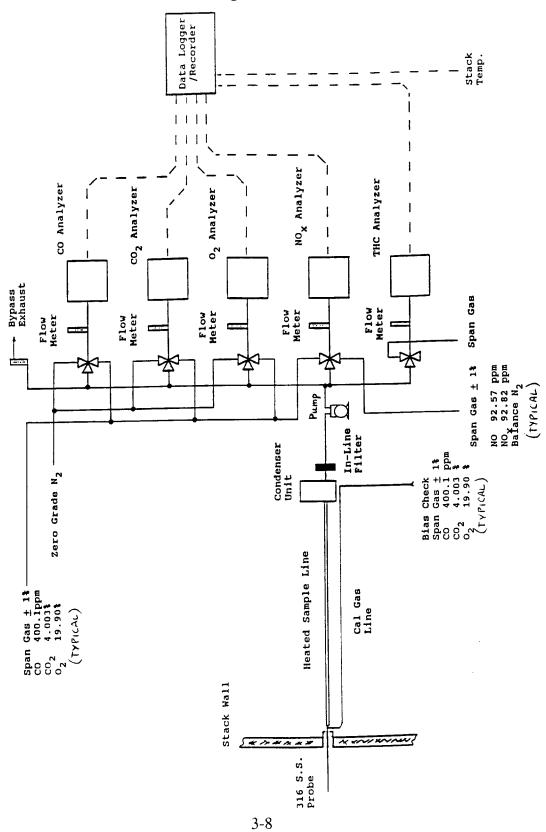
VOC Heated Hydrocarbon Analyzer Analyzer	- J.U.M. Engineeri	ng Model VE	-7 FID
Response Time	0-95% in less than	1.2 Seconds	
Zero Drift	± 1% of Full Sca	le in 24 Hrs	
Span Drift	+ 1% of Full scal	e in 24 Hrs	
Linearity	+ 1% of Full Sca	le	
Accuracy	Derived from the ± 1% gas was use		as
Sensitivity	1 ppb		
Range Change Consistency	Less than 1% Full	Scale	
Oxygen Synergism	Less than 2%		
Output	0-10.0 Vdc Scaled	l <b>:</b>	
	$egin{array}{c} R_1 \ R_2 \ R_3 \ R_4 \ R_5 \end{array}$	0-10 0-100 0-1,000 0-10,000 0-100,000	ppm as C <sub>3</sub>
Sample Flow Rate	3 Liters/Minute		

Table 3.2 Calibration Gases

Gas Composition	Use	Cylinder Ser. / No.	Certified Accuracy	Analysis Date
Nitrogen	Zero Gas	AAL / 4542	Zero Grade	N/A
11.0% O <sub>2</sub> 50.0 ppmv CO 11.0% CO <sub>2</sub> Bal N <sub>2</sub>	Span Gas	1L / 2572	± 1% ± 1% ± 1%	04/08/93
19.0% O <sub>2</sub> 75.0 ppm CO 18% CO <sub>2</sub> Bal N <sub>2</sub>	Span Gas	ALMO / 04292	<u>+</u> 1%	04/29/93
10.36 ppm NO 10.39 ppm NO <sub>x</sub> Bal N <sub>2</sub>	Span Gas	ALM / 010841	<u>+</u> 1%*	12/02/93 exp 12/95
22.90 ppm NO 23.46 ppm $NO_X$ Bal $N_2$	Span Gas	ALM / 034155	<u>+</u> 1%*	10/06/93 exp 10/95
43.83 ppm NO 44.17 ppm NO <sub>x</sub> Bal N <sub>2</sub>	Span Gas	ALM / 027046	<u>+</u> 1%*	12/23/93 exp 12/95
2.0% CO	Span Gas	ALM / 008830	<u>+</u> 1%*	09/17/93
4.0 CO	Span Gas	ALM / 033923	<u>+</u> 1%*	09/16/93

<sup>\*</sup> EPA Protocol 1 gas.

Figure 3.3 Continuous Monitoring Train - CARB Method 100



hydrocarbons, the instrument was used on a 0-50 ppmv range and standardized with 20 and 40 ppmv propane calibration gases. Each test run was about 50 minutes long.

In order to determine the methane concentration of the stack gas for correcting the total hydrocarbon monitoring data and as a back-up total hydrocarbon analysis, an integrated sample of the flue gas was collected for about 40 minutes concurrently with each monitoring run by using EPA Method 25 modified to eliminate the condensate trap. The sampling train is shown in Figure 3.4 and consisted of a stainless steel sampling probe connected through a flow control device (micro orifice disc meter) to an evacuated 12-liter stainless steel cylinder. The orifice meter reading and the cylinder gage vacuum were recorded at 10-minute intervals during the sampling period on field data sheets (see Appendix B).

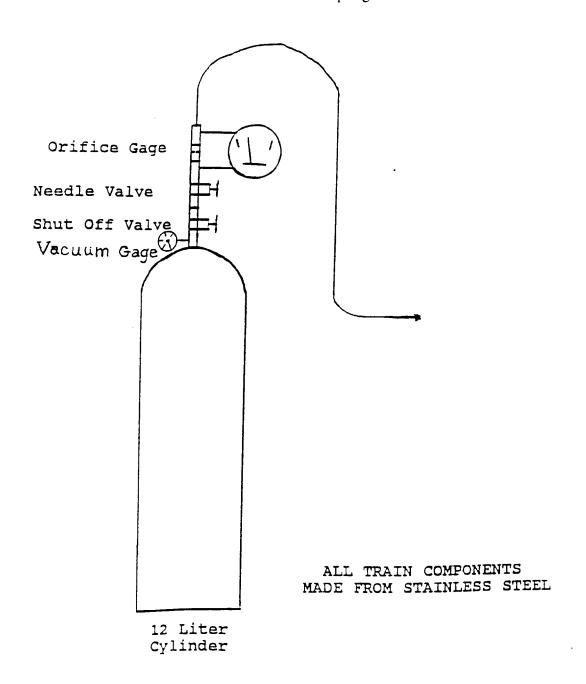
The integrated samples were analyzed for carbon monoxide (CO), methane  $(CH_4)$ , carbon dioxide  $(CO_2)$ , oxygen  $(O_2)$ , and nonmethane hydrocarbons (as  $C_1$ ) by Truesdail Laboratories, in Tustin, CA according to EPA Method 25. A sample submittal/chain of custody sheet was completed and is included in Appendix B along with the field data and calculation sheets. The analytical procedures performed at Truesdail are detailed in Appendix C along with the laboratory report.

Emission factors for the gases were calculated based on the natural gas usage. A mass balance calculation was performed on the entire power plant. The natural gas usage of all turbines is routed through a gas meter. The gas usage was monitored for first test day and related to the number of turbines operating and the kilowatt load for each.

#### Sulfur Dioxide

Sulfur dioxide emissions were expected to be very low. Since the source of sulfur is the odorant in the natural gas, the sulfur dioxide emissions were reported in pounds per therm (100 CF of natural gas) by using the sulfur content of the fuel (from the gas supplier).

Figure 3.4 VOC Sampling Train



NOTE: Samples usually collected in duplicate or triplicate concurrently

#### RESULTS

Calculations were made from the field data sheets and lab analyses to determine sample volume, molecular weight, velocities, flow rates, component concentrations, and micron cut-off size for the tests.

#### PM10

Analyses of PM10 emissions for each test run on the turbine included determinations for total solid particulate matter (PM). Condensables were not included in the PM10 results. Although some weight was recorded for each PM10 fraction, 1 milligram was chosen as a level of significance in order to account for fluctuations in the balance weights and for minute amounts of grease contamination from the threads of the PM10 sampling head. Results of the tests are summarized in Table 4.1. No significant PM10 or particulate matter was detected.

#### Gases

The nitrogen oxide and carbon monoxide emissions were 3.1 and 3.5 pounds per hour, respectively. The emission factors were reported as pounds per therm (100 CF) as indicated on the natural gas meter supplying the power plant. The gas meter had two indicators: one mechanical and one digital. The larger volume reading (digital) was assumed to be the corrected reading and this value was used for all of the emission factors calculations. However, this reading appears to have some unknown multiplier because the therm usage is too low for the known consumption of the gas turbines. For the purposes of calculating any periodic emissions, the emission factors should be used against the digital reading as indicated at the gas meter. Results are reported in Table 4.1.

The continuous monitoring for total hydrocarbons was unexpectedly variable and high (10 to 35 ppmv as  $C_3$ ) and indicated that the sample line, although heated, was contaminated from previous testing which involved very heavy organics. As a back-up, the integrated sample cylinders that were sampled concurrently and analyzed for methane were also analyzed for total nonmethane hydrocarbons and the resulting average value (5 ppmv) was used to calculate the VOC emission factor. Results are reported in Table 4.2.

The sulfur dioxide emission factor was calculated from information supplied by PG&E in San Francisco, CA which supplies the natural gas to the base. The sulfur dioxide emission factor used by PG&E is 0.001 pounds per million BTU which

Table 4.1

Turbine Number 2 Exhaust - PM10 Continuous Emission Monitoring

Description	Run #1	Run #2	Run #3	Average
Sampling Date Sample Number Turbine Load, KW Waste Heat Damper, % Gas Usage, therms/hr	12/03/93 OAFB-1 570 75 12.4*	12/03/93 OAFB-2 570 75 12.4*	12/03/93 OAFB-3 558 75 12.2*	
Flue Gas Temperature, °F Velocity, ft/sec Flow Rate, ACFM Flow Rate, DSCFM Moisture, % v/v	450 117.4 22,120 12,300 4.7	457 117.1 22,060 12,200 4.5	460 115.8 21,820 12,030 4.5	
PM10 Sample Start Sample Stop Sampling Time, min Sample volume, DSCF Cutoff, microns Collection, g Concentration, g/DSCF Emissions, lbs/hr Factor, lbs/therm	07:19 09:22 116.1 90.1 10.0 <0.001 <0.0002 <0.002 <0.002*	10:30 12:29 115.2 90.3 10.0 <0.001 <0.0002 <0.002 <0.002*	13:54 15:51 113.7 89.1 10.0 <0.001 <0.0002 <0.002 <0.002*	<0.0002 <0.02 <0.002
Total PM Collection, g Concentration, g/DSCF Emissions, lbs/hr	<0.001 <0.0004 <0.04	<0.001 <0.0004 <0.04	<0.001 <0.0004 <0.04	<0.0004 <0.04

Table 4.1 - Continued

Description	Run #1	Run #2	Run #3	Average
Gardina Manitorina Cosso				
Continuous Monitoring - Gases	09:56	11:51	13:46	
Sample Start			14:46	
Sample Stop	10:56	12:51	14:40	
Concentration, % v/v				
Carbon Dioxide	2.1	2.2	2.2	2.2
Oxygen	17.6	17.5	17.5	17.5
Concentration, ppmv				
Nitrogen Oxides	34	36	37	36
Carbon Monoxide	66	66	66	66
Emissions, lbs/hr				
Nitrogen Oxides	3.0	3.1	3.1	3.1
Carbon Monoxide	3.5	3.5	3.5	3.5
Factor, lbs/therm				1
Nitrogen Oxides	0.24*	0.25*	0.25*	0.25*
Carbon Monoxide	0.24	0.23*	0.29*	0.28*
Carbon Monoxide	0.20	0.20	0.27	0.20
	<u></u>			

<sup>\*</sup> Therms as indicated on digital readout of gas meter - appears to have some unknown multiplier.

Table 4.2
Turbine Number 2 Exhaust - Hydrocarbon Emissions

Description	Run #1	Run #2	Run #3	Average
Sampling Date Turbine Load, KW Waste Heat Damper, % Gas Usage, therms/hr	01/12/94 550 75 12.0*	01/12/94 550 75 12.0*	01/12/94 550 75 12.0*	
Flue Gas Temperature, °F Velocity, ft/sec Flow Rate, ACFM Flow Rate, DSCFM Moisture, % v/v	441 103.2 19,450 10,930 4.6**	444 103.2 19,450 10,930 4.6**	445 103.2 19,450 10,930 4.6**	
Hydrocarbon Samples Sample Start Sample Stop Sampling Time, min Concentration, % v/v Oxygen (O <sub>2</sub> ) Concentration, ppmv C <sub>1</sub> Carbon Dioxide (CO <sub>2</sub> ) Carbon Monoxide (CO) Methane (CH <sub>4</sub> ) Total HC (w/o CH <sub>4</sub> ) Emission Rate, lbs/hr C <sub>1</sub> Total HC (w/o CH <sub>4</sub> ) Emissions Factor, lbs/therm Total HC (w/o CH <sub>4</sub> )	10:56 11:36 40 17.9 18,610 60 7 <4 <0.04 <0.003*	12:18 12:58 40 18.2 18,590 58 6 11 0.22 0.018*	13:38 14:18 40 18.1 18,700 57 6 <4 <0.04 <0.003*	18.1 18,630 58 6 5

<sup>\*</sup> Therms as indicated on digital readout of gas meter - appears to have some unknown multiplier.

<sup>\*</sup> From PM10 testing - 12/3/93.

is equivalent to a concentration of 6 ppmv in the natural gas as hydrogen sulfide  $(H_2S)$ . Results are reported in Table 4.3.

Table 4.3

Turbine Number 2 Exhaust - Sulfur Dioxide Emission Factor

Sulfur Dioxide Emission Factor, lbs/therm	0.0001*
Emission Factor, 108/therm	0.0001

<sup>\*</sup> Based on standard emission factor used by gas supplier (PG&E).

## QUALITY ASSURANCE/QUALITY CONTROL

Source tests are performed to determine the types and amounts of pollutants emitted by a source. Information from this source test program may be used for obtaining permits, evaluating control equipment performance, updating emission inventories, and determining compliance with present and future emission regulations. For these purposes, reliable data are required. PES provides this reliability by using the following work practices:

#### USE OF STANDARD TEST PROCEDURES

CARB Methods 1 and 2 were utilized to measure flow rates. EPA Method 201A was used to determine the PM10 emission rate, and CARB Method 100 was used to determine the continuous emission rates for the gases. A procedure must be thoroughly studied under various conditions in order to be designated as a state or federal Method. Results of many executions of the procedure are compared to demonstrate accuracy and repeatability before adoption of the procedure as a source testing method.

#### USE OF TRAINED TEST PERSONNEL

Because of the complexity of typical source testing methods, testers should be trained and experienced with the test procedures in order to assure reliable results. PES personnel have had professional training and routinely conduct source tests.

### KNOWLEDGE OF SOURCE'S OPERATION

The source testing team should have sufficient knowledge of the process to be tested in order to properly document the process parameters during the tests. Without documentation of the process parameters used, results are much less meaningful. PES has previously tested boilers and combustion sources and is familiar with the processes and equipment.

## EQUIPMENT MAINTENANCE AND CALIBRATION

Use of properly maintained and calibrated test equipment is essential for minimizing systematic errors in results. All sampling devices were constructed,

maintained, and calibrated as suggested in EPA documents APTD-0576, and APTD-0581 (These are commonly accepted construction and maintenance manuals for source testing equipment). The dry gas meters were calibrated with a transfer gas meter with NBS traceability. These calibrations are included in Appendix D along with those for the nozzles, thermocouples, digital potentiometers, and Pitot tubes.

Quality control procedures used for continuous monitoring included the use of non-reactive 316 stainless steel or Teflon tubing and fittings throughout the system. A refrigeration unit was used with the pump down stream of the conditioned sample gas. All instrumentation was continuously monitored and checked between load conditions to insure data reliability during all sample runs. Bias checks were made with a calibration gas blend to confirm they met the tolerances specified in CARB Method 100.

All calibration gases were  $\pm 1$  per cent accuracy and provided by Scott Specialty Gases in San Bernardino, California. Copies of the calibration gas certifications are provided in Appendix D.

#### THOROUGH RECORD KEEPING

All data relating to the operation of the sampling train must be immediately recorded to ensure that it is not lost or misinterpreted. PES accomplishes this thorough record keeping by use of the field data sheets shown in Appendix B. The PES test team is familiar with these sheets and the information required to complete them. Any unusual occurrences in the process operation, unusual test instrument readings, or any other items that could affect the test results was also noted.

## USE OF THOROUGHLY CLEANED GLASSWARE

All glassware and probe lines were cleaned prior to the tests with hot tap water and then with 40 per cent nitric acid solution. The trains were then cleaned with 0.1 Normal sodium hydroxide solution, laboratory grade distilled water, air dried, and sealed until the tests.

# USE OF STANDARDIZED DATA REDUCTION TECHNIQUES

Data reduction was accomplished by the use of step by step calculation sheets. The calculations were systematic and easy to follow. All calculations for the source test are included in Appendix B.

# SUBMISSION OF BLANK SAMPLES

Filter and reagent samples from an unused but charged PM10 sampling train carried to the field was analyzed with the other samples to detect any possible contamination of sampling media or problems with lab analyses. No corrections were made to the measured concentrations of the collected samples, but the blank train results were reported on the calculation sheets.



# APPENDIX A EQUIPMENT PERMITS

— PACIFIC ENVIRONMENTAL SERVICES, INC. —



# BAY AREA AIR QUALITY MANAGEMENT DISTRICT

939 ELLIS STREET SAN FRANCISCO, CALIFORNIA 94109 (415) 771-6000

Plant# 232

Pauc.

Expires: **JUL 1, 1993** 

This document does not permit the holder to violate any District regulation or other tax

Gerald Reid Onizuka Air Force Base P O Box 3430 Sunnyvale, CA 94088 ORIGINAL SENT TO: Onizuka Air Force Base 1080 Lockheed Way Sunnyvale, CA 94088

S#	DESCRIPTION	[Schedule]	PAID
1	Turbine - Cogeneration, 12750K BTU/hr max, Multifue GAS TURBINE ENGINE WITH WASTE HEAT BOILER NO. 1 Emissions at: P1 Stack	1 [B]	150
2	Turbine - Cogeneration, 12750K BTU/hr max, Multifue GAS TURBINE ENGINE WITH WASTE HEAT BOILER NO. 2 Emissions at: P2 Stack	1 [B]	150
3	Turbine - Cogeneration, 12750K BTU/hr max, Multifue GAS TURBINE ENGINE WITH WASTE HEAT BOILER NO. 3 Emissions at: P3 Stack	1 [B]	150
4	Turbine - Cogeneration, 12750K BTU/hr max, Multifue GAS TURBINE ENGINE WITH WASTE HEAT BOILER NO. 4 Emissions at: P4 Stack	1 [B]	150
5	Turbine - Cogeneration, 12750K BTU/hr max, Multifue GAS TURBINE ENGINE WITH WASTE HEAT BOILER NO. 5 Emissions at: P5 Stack	1 [B]	150
6	Turbine - Cogeneration, 12750K BTU/hr max, Multifue GAS TURBINE ENGINE WITH WASTE HEAT BOILER NO. 6 Emissions at: P6 Stack	1 [B]	150
7	Turbine - Cogeneration, 12750K BTU/hr max, Multifue GAS TURBINE ENGINE WITH WASTE HEAT BOILER NO. 7 Emissions at: P7 Stack	1 · [B]	150
8	Turbine - Cogeneration, 12750K BTU/hr max, Multifue GAS TURBINE ENGINE WITH WASTE HEAT BOILER NO. 8 Emissions at: P8 Stack	1 [B]	150
9	Turbine - Cogeneration, 12750K BTU/hr max, Multifue GAS TURBINE ENGINE WITH WASTE HEAT BOILER NO. 9 Emissions at: P9 Stack	1 [B]	150



# APPENDIX B FIELD DATA AND CALCULATION SHEETS

PACIFIC ENVIRONMENTAL SERVICES, INC. —

# TRAVERSE POINT LOCATION FOR CIRCULAR DUCTS

DI ANT	ONIZUKA	AFB
PLANT	12-2-	93
SAMPLING LOCATION	TUKBIN	JE T
INSIDE OF FAR WALL TO OUTSIDE OF NIPPLE. (I	DISTANCE A)	25"
INSIDE OF NEAR WALL TO	)	j '/
OUTSIDE OF NIPPLE.	DISTANCE B)	24"
STACK I.D., (DISTANCE A- NEAREST UPSTREAM DIST	URBANCE	3 P
NEAREST DOWNSTREAM D	ISTURBANCE	2+ Ø
CALCULATOR		2110

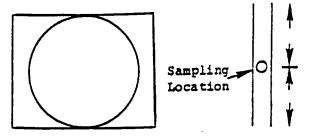
# SCHEMATIC OF SAMPLING LOCATION

TRAVERSE POINT NUMBER	FRACTION OF STACK I.D.	STACK I.D.	PRODUCT OF COLUMNS 2 AND 3 (TO NEAREST 1/8 INCH)	QISTANCE B	TRAVERSE POINT LOCATION FROM OUTSIDE OF NIPPLE (SUM OF COLUMNS 4 & 5)
		,			
	0.067	24	1.6		·
2	0.25	) )	6.0		
3 4	0.75	¥	18.0		1
4	0.93		22.3		
					·

- PACIFIC ENVIRONMENTAL SERVICES, INC. -

#### PRELIMINARY VELOCITY TRAVERSE

Plant:	MIZUK	A AFB	
Date:	12-	2-93	
Location	II TIR	BINE #2	)
Stack I.	D.:	<u> 2+"</u>	
Barometr	ic Pressure,	in. Hg:	30.3
	uge Pressure	, in. H <sub>2</sub> O:	
Operator	:s:		
	be I.D. Numb		SITA
Temperat	ure Readout	I.D.:	30X 3A
-	he Teak Chec		OK



Schematic of Traverse Point Layout

Traverse	Velo	city	Stack	Cyclo	nic	Traverse	Velo	oci ty	Stack	Cyclonic
Point	Head	(Δp <sub>s</sub> )	Temp.	Flow C	heck	Point	Head	$(\Delta p_g)$	Temp.	Flow Check
Number	in.	H <sub>2</sub> O	(Tq), °F	° from	Null	Number	in.	H <sub>2</sub> O	(Tg), °F	• from Null
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77 - CLASS LINER 17 A S.O. S.O. S.O. S.O. S.O. S.O. S.O. S.	Im- pinger Temp. °F 70 50 45 45	433					
S No. No.	Sample Box Temp. Filter Temp. °F	1 4 3 1 1					
i DiAMETER i DiAMETER i Bakerer Rate = CECC Leak Checl Leak Checl	Pump Vacuum In. IIg 4.0 4.0 7.0	444 44701					
angth and ube I.D. No. Holsture ox Number I.G. Dox Settl ce Dp st Leak B st Desat st Orsat	Inlet Outlet  Tmin e (Tmout)  59 54  71 57  72 61	69					
Probe Length Pitot Tube I. Nozzle I.D. P. Assumed Hoise Meter Box Num Heter Gamma Heter Gamma Heter Gamma Heter Camma Heter Camma Heter Tast Dr. Post Test Le. Post Test Pl. Post Test Pl. Post Test Pl. Post Test Pl.	Dry Gae   Inlet   (Tmin   F   71   71   72   72   72   72   72   72	36 36 36 76 7 7					
	Stack Temp. (Ta) °F 452 452 453	438 442 455 455 7-450					
FIELD DATA  THE STATE  TOP VIEW  Schematic of erse Point Layout  T. T. T. T. T. T. T. T. T. T. T. T. T. T	ce Pres. in. 1120 in. 1120 id Actus 1 2. 1 2. 1 2. 2 2. 2 2. 2 2. 2 2. 2	14.4.4.1   15.3     表					
FIELD DATA  THE STATES  TOP VIEW  Schematic of  Traverse Point Layout  Traverse Point Layout	Orifice Pres. Differential (All) in. 1120 Desired Actual  2.0 2.0 2.1 2.1 2.1 2.2 2.2 2.2	2.2 2.2 2.1 2.1 2.45					
1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	Velocity Head (APB) In. H20 2.7 2.7 2.5 2.5	2.2 2.2 2.2 2.2 30.2463 50.2463 50.2463 178					
20 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2	Gas Heter Reading (Vm) ft 423,473 435.0 447.0 447.0 457.1	48.4 482.5 503.1 503.1 513.852 113.852 448.21					
	/Clock / Timo /(24-hour clock) / 07:19 / 07:34 / 07:49 / 07:49	08:24 06:08 06:08 06:08 06:08 06:27 06:22					
g Location  Type  ber  Temperature  ric Pressure  Pressure (Ps)  Number(s)  t Leak Rate =  t Pitot Leak C  t Orsat Leak C  nd Record all	Sampling Time, (min) / (min) / 0 / 30 / 30 / 44.4 / 58.8 /	58.3 74.1 84.1 103.6 116.1 METER 073.3 29.40					
Sample Type  Run Humber Operator Ambient Temperature Barometric Pressure Static Pressure Filter Number(8) Pretest Leak Rate = 0.00 Pretest Orsat Leak Check Read and Record all Data	Traverse Point Number Rumber	(i) 22   FINI   FINI					
PACIFIC ENVIRONMENTAL SERVICES, INC.							



#### SAMPLE RETRIEVAL DATA

Plant:		DNIZ	UKA AFB	•
Date:			2-3-93	
Sampling Location	n:	π	IRBINE #2 6	STACK
Sampling Type (M	ethod):		PM-10	
Run Number: 3			BAB DAY	B-1
Sample Box Numbe	r:		38	
Clean-up Man:			BROWN, KEAR	NEY
			F028	
Comments:				
Filter				
Filter Number: Description of F	ilter: <u>CUE</u>	łN		
Moisture	<b>#</b> /		#2-	<b>*</b> 3
Impingers: Final Volume: Initial Volume: Net Volume: Total H <sub>2</sub> O:	157 0 100.0 57.0 57	mL mL mL mL	102,0 m 100.0 m 2.0 m	L <u>0.0</u> mI L 16.0 mI
Silica Gel				
Final Volume: Initial Volume: Net Volume: Total Moisture:	700,9 681-9 14.0 94	a a a a	a a a a	a a a
Description of Ir	npinger catch	:	CLEAR	

PACIFIC ENVIRONMENTAL SERVICES, INC. —

Date: 12-3-93

Source/Sample Number: OAFB-1

- 1.  $Vm(std) = (17.64)(Vm)(Y) \left[ \frac{P_{bar} + (\Delta H/13.6)}{Tm} \right]$   $Vm(std) = (17.64)(89.879)(0.99) \left[ \frac{(30.2) + (2.15/13.6)}{(52.9)} \right]$   $Vm(std) = \frac{90.1}{} dscf.$
- 2. Volume water vapor collected (standard conditions).

V(1o) = 94 condensate from impingers and selica gel.

$$Vw(std) = (0.04707) V(10) = (0.04707) (9 \%)$$

Vw(std) = Y.42 scf.

3. Percent moisture, by volume.

$$Bw_{S} = \frac{Vw(std)}{Vw(std) + Vm(std)} = \frac{(4.42)}{(4.42) + (90.1)} = \frac{0.047}{}$$

 $BW_{S} = 4.7/.$ 

4. Molecular weight, stack gas.

Dry molecular weight.

$$Md = 0.440(\% CO_2) + 0.320(\% O_2) + 0.280(\% N_2 + \% CO)$$

$$Md = 0.440 (2.5) + 0.320(17.5) + 0.280(80)$$

 $Md = 29.1 \quad 1b/1b-mole.$ 

$$Ms = Md + Bw_s (18 - Md) = (29.1) + (0.647)(18 - 29.1)$$

Ms = 28.58 lb/lb-mole.

Date: 12-3-93

Source/Sample Number: <u>&AFB-1</u>

5. Stack gas velocity average.

$$Vs(avg) = (85.49)(Cp)(\sqrt{\Delta P}) \left[ avg \sqrt{\frac{(Ts)}{(Ps)(Ms)}} \right]$$

$$Vs(avg) = (85.49)(0.84)(1.59) \sqrt{\frac{(910)}{(30.1)(28.58)}}$$

Vs(avg) = 117.4 ft/sec.

6. Stack volumetric flow rate, actual conditions (stack temperature and pressure).

$$Qs = (60)(Vs)(A) = (60)(117.4)(3.14)$$

Qs = 22118 acfm.

7. Stack volumetric flow rate, standard conditions (68 degrees F, 29.92 Hg).

$$Q(std) = (17.64)(Qs)(1-Bw_s) \frac{(Ps)}{(Ts)}$$

$$Q(std) = (17.64)(22/18)(1 - 0.847) \begin{bmatrix} 30.1 \\ 9/0 \end{bmatrix}$$

 $Q(std) = 12299 \quad dscfm.$ 

8. Isokinetic variation.

$$ZI = (K) \left[ \frac{(Ts)(Vm(std))}{(Ps)(Vs)(An)(\theta)(1 - Bw_s)} \right]$$

$$3I = (0.0945)$$
 ( )( )( )(1 - )



Date: 12-3-73

9. Viscosity of stack gas:

 $\mu_s = 152.418 + 0.2552 t_s + 3.2355x10^{-5} (t_s)^2 +$ 

0.53147 (%O<sub>2</sub>) - 74.143 Bw<sub>s</sub>

 $\mu_s = 152.418 + 0.2552(450) + 3.2355 \times 10^{-5} (450)^2 +$ 

0.53147(17.5) - 74.143(0047)

 $\mu_s = 279.0$  micropoise.

10. Cyclone flow rate:

 $Q_s = 0.002837 \ \mu_s \ \left[ \frac{t_s + 460}{M_s \ P_s} \right]^{0.2949}$ 

 $Q_s = 0.002837 (279.6) \left( \frac{(450) + 460}{(23.58) (30.1)} \right)^{0.2949}$ 

 $Q_{a} = 0.81$  ft<sup>3</sup>/min

11. Orifice pressure head for cyclone flow rate:

 $\Delta H = \left( \frac{Q_s (1 - Bw_s) P_s}{t_s + 460} \right)^2 \left( \frac{(t_m + 460) M_d (1.083) \Delta H@}{P_{bar}} \right)$ 

 $\Delta H = \left( \begin{array}{c} (0.0) \ ) \ (1-0.047) \ (30.1) \end{array} \right)^{2} \ \left( \begin{array}{c} (67 \ + 460) \ (29.1) \ (1.083) \ (1.916) \end{array} \right)$ 

 $\Delta H = 0.69$  in.  $H_2O$ 



Date: 12-3-93

Source/Sample Number: OAFB-1

Stack viscosity,  $\mu_s$ , micropoise =  $\frac{279.6}{30.1}$ Absolute stack pressure,  $P_s$ , in. Hg =  $\frac{30.1}{450}$ Average stack temperature,  $t_s$ ,  $F = \frac{450}{69}$ Method 201A pitot coefficient,  $C_p = \frac{69}{0.84}$ Cyclone flow rate, ft min,  $Q_s = \frac{0.81}{0.34}$ Method 2 pitot coefficient,  $C_s = \frac{0.81}{0.34}$ Molecular weight of stack gas, wet basis,  $M_s = \frac{28.58}{0.150}$ Nozzle diameter,  $D_s$ , in. =  $\frac{0.150}{0.150}$ 

Nozzle velocity

$$v_n = \frac{3.056 \, Q_s}{D_n^2} = \frac{110.0}{10.0} \, \text{ft/sec}$$

Maximum and minimum velocities:

Calculate  $R_{min}$ .

If  $R_{min}$  is less than 0.5, or if an imaginary number occurs when calculating  $R_{min}$ , use Equation 1 to calculate  $v_{min}$ . Otherwise, use Equation 2.

Eq. 1 
$$v_{min} = v_n (0.5) = ____ ft/sec$$
  
Eq. 2  $v_{min} = v_n R_{min} = _{82.5} ft/sec$ 

Calculate  $R_{\text{max}}$ .

$$R_{\text{max}} = 0.4457 + \sqrt{0.5690 + \frac{0.2603 (\sqrt{Q_s}) \mu_s}{v_n^{1.5}}} = 1.24$$



Plant: ONIZUKA AFB

Date: 12-3-93

Source/Sample Number: OAFB-1

If  $R_{max}$  is greater than 1.5, use Equation 3 to calculate  $v_{max}$ .

Eq. 3 
$$v_{max} = v_n (1.5) = ____ ft/sec$$
  
Eq. 4  $v_{max} = v_n R_{max} = _136.4$  ft/sec

Maximum and minimum velocity head values:

$$\Delta p_{min} = 1.3686 \times 10^{-4} \frac{P_s M_w (v_{min})^2}{(t_s + 460) C_p^2} = \frac{1.25}{10.00} in. H_20$$

$$\Delta p_{\text{max}} = 1.3686 \times 10^{-4} \frac{P_s M_w (v_{\text{max}})^2}{(t_s + 460) C_p^2} = \frac{3.41}{\text{in. H}_20}$$

Calculate the actual  $D_{50}$  of the cyclone for the given conditions as follows:

$$D_{50} = \beta_1 \left( \frac{t_s + 460}{M_w P_s} \right)^{0.2091} x \left( \frac{\mu_s}{Q_s} \right)^{0.7091}$$

where,  $\beta_1 = 0.15625$ 

$$D_{50} = (0.15625) \left( \frac{910}{(2858)(30.1)} \right)^{0.2091} \times \left( \frac{279.6}{0.81} \right)^{0.7091}$$

$$D_{50} = 9.97 \mu m$$



CLIENT: ONIZUKA AFB

Project No. F028

PM 10 TEST #1

# PARTICULATE CALCULATIONS

	TURRU	NE #2 EXHAUST	- Most Date:	12-03-93
Sample Numbe	r: <u>OAFB-1</u>	Sample Vo	lume: <u>40./</u>	DSCF
Stack Flow R	ate: <u>/2,300</u>	DSCFM	·	
<u>Particulate</u>	Catch: (gr	ams) PMIO	- PM10	
Filter:	Final Weigh Initial Weigh Net Weigh	t <u>0.2084</u> t <u>0.2083</u> t <u>0.000</u>		
ACETONE I <del>mpinger:</del>	Final Weigh Initial Weigh Net Weigh	t $\frac{28.6871}{28.6878}$ t $\frac{-0.0007}{}$	29.0835 29.0832 0.0003	
Extract:	Final Weigh Initial Weigh Net Weigh	t		
Total:		<u>&lt;0.0010</u>	40.0010	
<u>Particulate</u>	Concentration:			
( 40.00)	( 90./ )	43 grains/gram DSCF	= <0.0002	grains/DSCF
Particulate	Emissions:			
(20.0004 g)	r/DSCF x ( /2,30 7000 gr	00 ) DSCFM x 60 ains/lb	<u> min/hr</u> = _ <	0.02 lbs/hr
Rule 404 Lim	itation @^	<u> </u>	grains	/DSCF
Rule 405 Lim	itation @	1bs/hr = _	lbs/	hr

— PACIFIC ENVIRONMENTAL SERVICES, INC. —

3			
ر ا الال	NIN. 15.00 1	S. 7. 7. 7. 7. 7. 7. 7. 7. 7. 7. 7. 7. 7.	
5 FT - 6 (145) (1 (145))  - 17 A  - 18 A  - 18	Im- plnyer Temp. %	53	9 9
Ky   M   N   N   N   N   N   N   N   N   N	Sample Box Temp. Filter Temp. °F	1 1 1 1	
	Pump Vacuum In. Hg	1,800	
ngth and be I.D. No. Hoisture K Number	Dry Gas Meter Temp. Inlet Outlet Tmin * (Tmout) * F 74 70 83 74 70 83 74 70 83 74 70	85. 89. 89. 87. 87.	
Probe Length Pitot Tube I. Nozzle I.D. n Assumed Noise Assumed Noise Reter Box Nur Heter Alle C Factor Heter Gamma	Dry Gas H Inlet (Tmin) *P	95 100 98 96 96 12	
ayout 14.EK	Stack Temp. (Tg) 	464	15.0-21
FIELD DATA  (SEE TEST 1)  Schematic of erse Point Layout  Incharek	Pres. 120 Actual 2.7 2.7 2.3 2.3	richie 2	\$\frac{\pi}{2}
FIELD DATA  (SEE TEST 1)  Schematic of Traverse Point Layout Traverse Point Layout	Orifice Pres. Differential (All In. 1120 Desired Actual 2.2. 2.2. 2.3. 2.3. 2.3. 2.3. 2.3.	2.3 2.3 2.3 2.3 2.3 04:	३.4.६
1 1 4 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	Velocity Head (DPg) In. H20 2.7 2.7 2.7 2.7 2.2	27.25 2.8 27.25 2.8	1.59 302641 32495 264
2 12 2 2 2	Gas Meter Reading (Vm.) ft 513.919 524.3 536.4 547.7 553.3	558.8 571.6 534.3 595.7 607.011	498 363 998 707 344
CLCS Check Check Check	/Clock / Timo /(24-hour clock) / 10: 30 / 10: 45 / 11: 06 / 11: 14	11:31 11:46 12:01 12:24 12:24	0018
ocation  e Pressure P	Sampling Time, (min) (min) (0 15.0 36.0 43.5 43.5	57.0 7 <b>6.</b> 3 8 <b>5.</b> 0 101.1 115.2	Me 18K
Plant CNNZU  Date 1.2  Sampling Location  Sample Type  Run Number  Operator  Ambient Temperature  Barometric Pressure  Static Pressure (Pg Filter Number(s)  Pretest Leak Rate =  Pretest Orsat Leak  Read and Record all	Traverse Point Numbor L + 2 2 (1) 2 FINI	5 E E E E E E E E E E E E E E E E E E E	eA5
	PACIFIC ENVIRCINMENTAL SERVICES	s. Inc	



## SAMPLE RETRIEVAL DATA

Plant:	ON	IIZUKA	AFB				
Date: 12-3-93							
Sampling Location: _		TURBIN	E#2 STACK				
Sampling Type (Metho	d):	P	M-10				
Run Number:		C	AFB-2				
Sample Box Number:			3B				
Clean-up Man: BROWN KEARNEY							
Job Number:			028				
Comments:			020				
Commencs:							
		,					
Filter							
* 11261		•					
Filter Number:							
Description of Filte	r: CLEA	FN -			_		
Descripcion or 11100		<del>``</del>					
Moisture							
MOISCUIE	#1		#2	<b>*</b> 3			
	<b>–</b> (		# C	F 3			
Impingers:							
Final Volume:	156.0	mL	108mL	3.0m	αL		
Initial Volume:	100.0	mL	100.0 mL		nL		
<del>-</del>	56	mil.	7 mL		αL		
Net Volume:	56	mL	64 mL		nL		
Total H <sub>2</sub> O:	56				_		
		:					
		í					
Silica Gel		r -					
	682.3	~	g		ĭ		
Final Volume:	659.7	——;ā -					
Initial Volume:		a -	a				
Net Volume:	22.6	a -	g				
Total Moisture:	\$9.6	a -	g		j		
		1					
	4-1-	. ^	1 CAO				
Description of Inpir	ger catch	:	LEAR				
		<del></del>		···	_		
		<del></del>					
		Ţ					

PACIFIC ENVIRON MENTAL SERVICES, INC.

Date: 12-3-93

Source/Sample Number: <u>OAFB-2</u>

- 1.  $Vm(std) = (17.64)(Vm)(Y) \begin{bmatrix} P_{bar} + (\Delta H/13.6) \\ Tm \end{bmatrix}$   $Vm(std) = (17.64)(93.692)(6.99) \begin{bmatrix} (36.2) + (2.3)/13.6 \\ (547) \end{bmatrix}$   $Vm(std) = 90.3 \quad dscf.$
- 2. Volume water vapor collected (standard conditions).

V(10) = 89.6 condensate from impingers and selica gel.

Vw(std) = (0.04707) V(10) = (0.04707)(89.6)

Vw(std) = 4.22 scf.

3. Percent moisture, by volume.

 $BW_S = \frac{VW(std)}{VW(std) + Vm(std)} = \frac{(4.22)}{(4.22) + (90.3)} = \frac{0.045}{}$ 

 $Bw_s = 4.5 i$ .

4. Molecular weight, stack gas.

Dry molecular weight.

 $Md = 0.440(\% CO_2) + 0.320(\% O_2) + 0.280(\% N_2 + \% CO)$ 

Md = 0.440 (2.5) + 0.320(7.5) + 0.280(80)

Md = 29.1 1b/1b-mole.

 $Ms = Md + Bw_s (18 - Md) = (29.1) + (0.045)(18 - 29.1)$ 

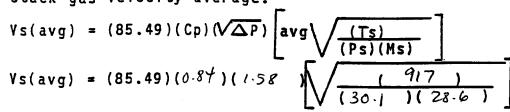
Ms = 78.6 lb/lb-mole.



Date: 12-3-93

Source/Sample Number: <u>OAFB-2</u>

5. Stack gas velocity average.



Vs(avg) = 117.1 ft/sec.

6. Stack volumetric flow rate, actual conditions (stack temperature and pressure).

$$Qs = (60)(Vs)(A) = (60)(117.()(3.) \neq )$$

Qs = 22062 acfm.

7. Stack volumetric flow rate, standard conditions (68 degrees F, 29.92 Hg).

$$Q(std) = (17.64)(Qs)(1-Bw_s)[(Ps)]$$

$$Q(std) = (17.64)(22062)(1 - 0.845) \begin{bmatrix} 301 \\ 917 \end{bmatrix}$$

Q(std) = 12200 dscfm.

8. Isokinetic variation.

$$%I = (K) \left[ \frac{(Ts)(Vm(std))}{(Ps)(Vs)(An)(\theta)(1 - Bw_s)} \right]$$

$$z_{I} = (0.0945) \left[ \frac{()())}{()()()()()()} \right]$$



Date: 12-3-93

Source/Sample Number: OAFB-2

9. Viscosity of stack gas:

$$\mu_s = 152.418 + 0.2552 t_s + 3.2355 \times 10^{-5} (t_s)^2 +$$

0.53147 (%O<sub>2</sub>) - 74.143 Bw<sub>s</sub>

$$\mu_s = 152.418 + 0.2552(457) + 3.2355 \times 10^{-5} (457)^2 +$$

0.53147 (17.5) - 74.143 (0.045)

 $\mu_{*} = \frac{781.8}{}$  micropoise.

10. Cyclone flow rate:

$$Q_s = 0.002837 \ \mu_s \ \left(\frac{t_s + 460}{M_s \ P_s}\right)^{0.2949}$$

$$Q_{s} = 0.002837 (281.8) \left( \frac{(3757) + 460}{(28.6)(30.1)} \right)^{0.2949}$$

 $Q_s = 0.81$  ft<sup>3</sup>/min

11. Orifice pressure head for cyclone flow rate:

$$\Delta H = \left( \frac{Q_s (1-Bw_s) P_s}{t_s + 460} \right)^2 \left( \frac{(t_m + 460) M_d (1.083) \Delta H@}{P_{bar}} \right)$$

$$\Delta H = \left| \frac{(0.81)(1-0.045)(301)}{(457) + 460} \right|^{2} \left| \frac{(87 + 460)(291)(1.083)(1.916)}{(30.2)} \right|$$

 $\Delta H = \delta \cdot 71$  in.  $H_2O$ 



Date: 12-3-93

Source/Sample Number: <u>OAFB-2</u>

Stack viscosity,  $\mu_s$ , micropoise =  $\frac{281.8}{30.i}$ Absolute stack pressure,  $P_s$ , in. Hg =  $\frac{30.i}{457}$ Meter temperature,  $t_s$ , \*F =  $\frac{457}{57}$ Method 201A pitot coefficient,  $C_s$  =  $\frac{0.84}{57}$ Method 2 pitot coefficient,  $C_s$  =  $\frac{0.84}{57}$ Method 2 pitot coefficient,  $C_s$  =  $\frac{0.84}{57}$ Molecular weight of stack gas, wet basis,  $M_s$  =  $\frac{0.83}{50}$ Nozzle diameter,  $D_s$ , in. =  $\frac{0.150}{50}$ 

Nozzle velocity

$$v_n = \frac{3.056 \, Q_s}{D_n^2} = \frac{110.0 \, \text{ft/sec}}{}$$

Maximum and minimum velocities:

Calculate R<sub>min</sub>.

If  $R_{\min}$  is less than 0.5, or if an imaginary number occurs when calculating  $R_{\min}$ , use Equation 1 to calculate  $v_{\min}$ . Otherwise, use Equation 2.

Eq. 1 
$$v_{min} = v_n (0.5) =$$
\_\_\_\_ ft/sec  
Eq. 2  $v_{min} = v_n R_{min} =$ \_82.5 ft/sec  
Calculate  $R_{max}$ .

$$R_{\text{max}} = 0.4457 + \sqrt{0.5690 + \frac{0.2603 \left(\sqrt{Q_s}\right) \mu_s}{v_n^{1.5}}} = 1.24$$



Date: 12-3-93

Source/Sample Number: <u>6AFB-2</u>

If  $R_{\text{max}}$  is greater than 1.5, use Equation 3 to calculate  $v_{\text{max}}$  . Otherwise use Equation 4.

Eq. 3 
$$v_{max} = v_n (1.5) = ____ ft/sec$$
  
Eq. 4  $v_{max} = v_n R_{max} = ___ /36.4$  ft/sec

Maximum and minimum velocity head values:

$$\Delta p_{min} = 1.3686 \times 10^{-4} \frac{P_s M_w (v_{min})^2}{(t_s + 460) C_p^2} = 1.27$$
 in. H<sub>2</sub>0

$$\Delta p_{\text{max}} = 1.3686 \times 10^{-4} \frac{P_s M_w (v_{\text{max}})^2}{(t_s + 460) C_p^2} = 3.39 \text{ in. } H_20$$

Calculate the actual  $D_{50}$  of the cyclone for the given conditions as follows:

$$D_{50} = \beta_1 \left( \frac{t_s + 460}{M_w P_s} \right)^{0.2091} \times \left( \frac{\mu_s}{Q_s} \right)^{0.7091}$$

where,  $\beta_1 = 0.15625$ 

$$D_{50} = (0.15625) \left( \frac{917}{(28 \cdot 6)(36)} \right)^{0.2091} \times \left( \frac{281.8}{0.81} \right)^{0.7091}$$

$$D_{50} = 16.5 \% \mu m$$



CLIENT: ONIZUKA AFB
Project No. F028

	PART	CULAT	E CALCULAT	ions /	MIO	TEST #2
Sampling Loc	ation:	BINE	#2 EXHAUST	Test D	ate: <u>/</u>	2-03-94
Sample Number			Sample Vol	ume:	90.3	DSCF
Stack Flow R	ate: 12,20	0 DS	CFM			
Particulate (	Catch: (	grams)	PMIO	> PMIC		
Filter:	Final Weight Initial Weight Weight Weight Weight Final Weight Weight Weight Final Weight Weight Weight Final Weight Weigh	ght	0.2012			
ACETONE Impinger:	Final Weig Initial Weig Net Weig	ght <u>2</u> ght <u>2</u>	8.6117 8.6117	28.34 28.34 0.00	56 43 13	
Extract:	Final Weig Initial Weig Net Weig	ght _				
Total:			0.0010	0.001	<u>3</u> _	
Particulate Concentration:						
Particulate Emissions:						
(20,0002) gr/DSCF x $(12,200)$ DSCFM x 60 min/hr = $20.02$ lbs/hr 7000 grains/lb						
Rule 404 Lim	itation @	NA	SCFM =	g:	rains/D	scF
Rule 405 Lim	itation @	NA	lbs/hr = _		lbs/hr	

PACIFIC ENVIRONMENTAL SERVICES, INC. -



#### SAMPLE RETRIEVAL DATA

Plant:(	DNIZUKA AFB		
Date:	12-3-93	) 	
Sampling Location:	TURBINE !	2 STACK	
Sampling Type (Method):	PM-10	<u> </u>	
Run Number:	OAFB	- 3	
Sample Box Number:	3B		
Clean-up Man:	* KOWN	KEARNEY	
Job Number:		8'	
Comments:			
Filter			
Filter Number:			
Description of Filter:			
Moisture			
Impingers: Final Volume: 154.	n	106 .DmL	#3 <u>0</u> mL
		$\frac{100 \cdot 0}{100 \cdot 0} \text{ mL}$	$\frac{700}{0.0}$ mL
		6.0 mL	3,0 mL
	mL	60 mL	63 mL
Total $H_2O$ :		<u> </u>	
Silica Gel			
	. 2 g	g	g
Initial Volume: 690		g	a
Net Volume: 25		g	g
Total Moisture: % %.	<u> </u>	g	a
Description of Inpinger cat	ch:		

PACIFIC ENVIRONMENTAL SERVICES, INC.

Date: 12-3-93

Source/Sample Number:\_\_\_OAFB-3

- 1.  $Vm(std) = (17.64)(Vm)(Y) \left[ \frac{P_{bar} + (\Delta H/13.6)}{Tm} \right]$   $Vm(std) = (17.64)(91.077)(0.99) \left[ \frac{(30.2) + (2.2)(13.6)}{(542)} \right]$   $Vm(std) = 89-1 \quad dscf.$
- 2. Volume water vapor collected (standard conditions).

V(10) = 88.1 condensate from impingers and selica gel.

Vw(std) = (0.04707) V(10) = (0.04707)(884)

Vw(std) = 4-15 scf.

3. Percent moisture, by volume.

 $Bw_{s} = \frac{Vw(std)}{Vw(std) + Vm(std)} = \frac{(4.15)}{(4.15) + (89.1)} = \frac{0.045}{(4.15)}$ 

 $BW_S = \underline{4.51}.$ 

4. Molecular weight, stack gas.

Dry molecular weight.

 $Md = 0.440(\% CO_2) + 0.320(\% O_2) + 0.280(\% N_2 + \% CO)$ 

Md = 0.440 (2.5) + 0.320(17.5) + 0.280(80)

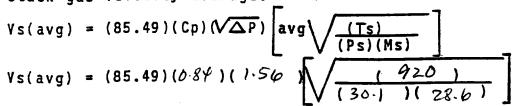
 $Ms = Md + Bw_s (18 - Md) = (29.1) + (0.645)(18 - 29.1)$ 

Ms = 28.0 lb/lb-mole.

Date: 12-3-93

Source/Sample Number: OAFB-3

5. Stack gas velocity average.



Vs(avg) = 115.8 ft/sec.

6. Stack volumetric flow rate, actual conditions (stack temperature and pressure).

$$Qs = (60)(Vs)(A) = (60)(115.8)(3.14)$$

Qs = 21817 acfm.

7. Stack volumetric flow rate, standard conditions (68 degrees F, 29.92 Hg).

$$Q(std) = (17.64)(Qs)(1-Bw_s)[(Ps)]$$

$$Q(std) = (17.64)(21817)(1 - 0.045) [30.1]$$

Q(std) = 1202-5 dscfm.

8. Isokinetic variation.

$$2I = (K) \left[ \frac{(Ts)(Vm(std))}{(Ps)(Vs)(An)(\theta)(1 - Bw_s)} \right]$$



Date: 12-3-93

Source/Sample Number: OAFB-3

### 9. Viscosity of stack gas:

$$\mu_s = 152.418 + 0.2552 t_s + 3.2355x10^{-5} (t_s)^2 +$$

$$\mu_{\rm s} = 152.418 + 0.2552(460) + 3.2355 \times 10^{-5} (460)^2 +$$

$$0.53147(17.5) - 74.143(0.045)$$

 $\mu_s = 282.6$  micropoise.

### 10. Cyclone flow rate:

$$Q_s = 0.002837 \ \mu_s \ \left| \frac{t_s + 460}{M_s P_s} \right|^{0.2949}$$

$$Q_{s} = 0.002837(282.6) \left( \frac{(460) + 460}{(28.6)(30)} \right)^{0.2949}$$

$$Q_{\bullet} = 0.82$$
 ft<sup>3</sup>/min

## 11. Orifice pressure head for cyclone flow rate:

$$\Delta H = \left( \frac{Q_s (1-Bw_s) P_s}{t_a + 460} \right)^2 \left( \frac{(t_m + 460) M_d (1.083) \Delta H@}{P_{bar}} \right)$$

$$\Delta H = \left| \frac{(0.82)(1-0.045)(30.1)}{(460) + 460} \right|^2 \left| \frac{(82 + 460)(29-1)(1.083)(1.916)}{(30.2)} \right|$$

$$\Delta H = 0.71$$
 in.  $H_2O$ 



Date: 12-3-93

Source/Sample Number: <u>OAFB-3</u>

Stack viscosity,  $\mu_s$ , micropoise = 282.6Absolute stack pressure,  $P_s$ , in. Hg = 30.1Average stack temperature,  $t_s$ , F = 460Meter temperature,  $t_m$ , F = 82Method 201A pitot coefficient,  $C_s = 0.84$ Cyclone flow rate,  $ft^3$ /min,  $Q_s = 0.84$ Method 2 pitot coefficient,  $C_s = 0.84$ Molecular weight of stack gas, wet basis,  $M_s = 28.6$ Nozzle diameter,  $D_s$ , in. = 0.150

Nozzle velocity

$$v_n = \frac{3.056 \, Q_s}{D_n^2} = \frac{11.4}{\text{ft/sec}}$$

Maximum and minimum velocities:

Calculate R<sub>min</sub>.

If  $R_{\min}$  is less than 0.5, or if an imaginary number occurs when calculating  $R_{\min}$ , use Equation 1 to calculate  $v_{\min}$ . Otherwise, use Equation 2.

Eq. 1 
$$v_{min} = v_n (0.5) = ____ ft/sec$$
  
Eq. 2  $v_{min} = v_n R_{min} = _83.66 ft/sec$ 

Calculate  $R_{max}$ .

$$R_{\text{max}} = 0.4457 + \sqrt{0.5690 + \frac{0.2603 \left(\sqrt{Q_s}\right) \mu_s}{v_n^{1.5}}} = 1.24$$



Date: 12-3-93

Source/Sample Number: <u>OAFB-3</u>

If  $R_{\text{max}}$  is greater than 1.5, use Equation 3 to calculate  $v_{\text{max}}.$  Otherwise, use Equation 4.

Eq. 3 
$$v_{max} = v_n (1.5) = ____ ft/sec$$
  
Eq. 4  $v_{max} = v_n R_{max} = _{138.1} ft/sec$ 

Maximum and minimum velocity head values:

$$\Delta p_{min} = 1.3686 \times 10^{-4} \frac{P_s M_w (v_{min})^2}{(t_s + 460) C_p^2} = 1.27 \text{ in. } H_20$$

$$\Delta p_{\text{max}} = 1.3686 \times 10^{-4} \frac{P_s M_w (v_{\text{max}})^2}{(t_s + 460) C_p^2} = 3.40 \text{ in. } H_20$$

Calculate the actual  $D_{50}$  of the cyclone for the given conditions as follows:

$$D_{50} = \beta_1 \left( \frac{t_s + 460}{M_w P_s} \right)^{0.2091} x \left( \frac{\mu_s}{Q_s} \right)^{0.7091}$$

where,  $\beta_1 = 0.15625$ 

$$D_{50} = (0.15625) \left( \frac{920}{(28 \cdot 6)(30 \cdot i)} \right)^{0.2091} \times \left( \frac{282 \cdot 6}{0.32} \right)^{0.7091}$$

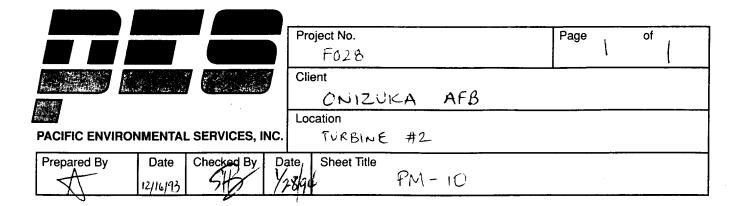
$$D_{50} = 9.98 \mu m$$



Project No. F028

			•			
	<u>PARTICU</u>	LATE CALCULATI	IONE PMIO	TEST #3		
Sampling Location:						
Sample Number:			ume: 89.1	DSCF		
Stack Flow Rate:	12,030	DSCFM	·			
Particulate Catch:	(gram	s) PMIO	> PMIO			
Filter: Fina Initia Ne	l Weight l Weight t Weight	0.2056 0.2065 - 0.0009				
	l Weight	28.6882 28.6882 0.0000	28.1727 28.172/ 0.0006			
Initia	l Weight l Weight : Weight					
Total:		20.0010	0,0006			
Particulate Concent						
(20.00/0) grams	9,1 ) D	grains/gram SCF	= <0.0002	grains/DSCF		
Particulate Emissions:						
( <sup>20,000</sup> 2) gr/DSCF x	//2,030 000 grain	) DSCFM x 60 p	$min/hr = \angle 0$	02 lbs/hr		
Rule 404 Limitation	e NA	SCFM =	grains/	DSCF		
Rule 405 Limitation	e NA	lbs/hr =	lbs/h	ır		

-- PACIFIC ENVIRONMENTAL SERVICES, INC. --



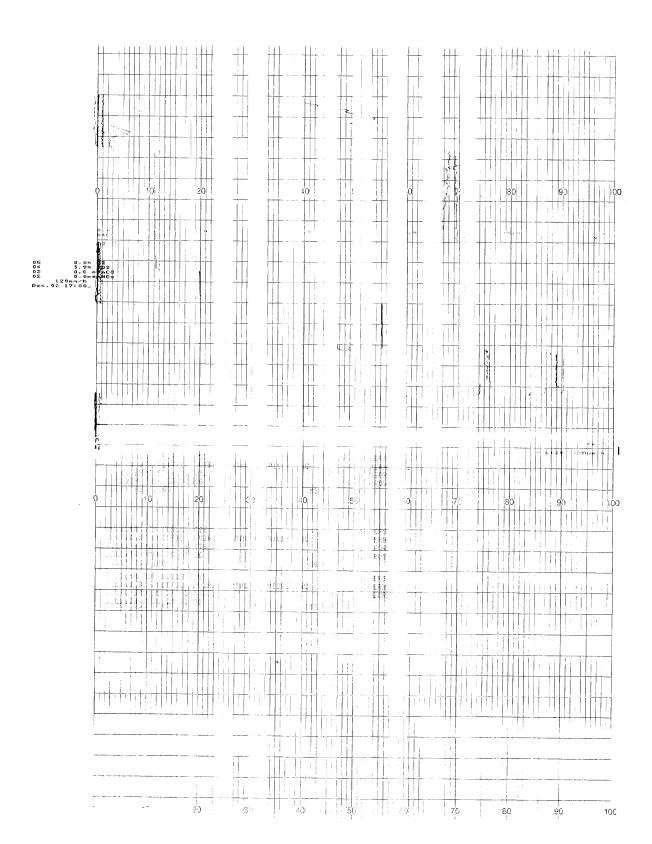
BEAKER	POST WEIGHT &	PRE WEIGHT 9	NET g
IA	29.0035	29.5832	0.0003
2A	28.3456	28.3443	0.0013
3A	28.1727	28.1721	0.0006
18	28.6871	28. 6£78	-0.0007
28	28.6117	28.6117	0. 0000
3B	28.6882	23. 5382	0.0000

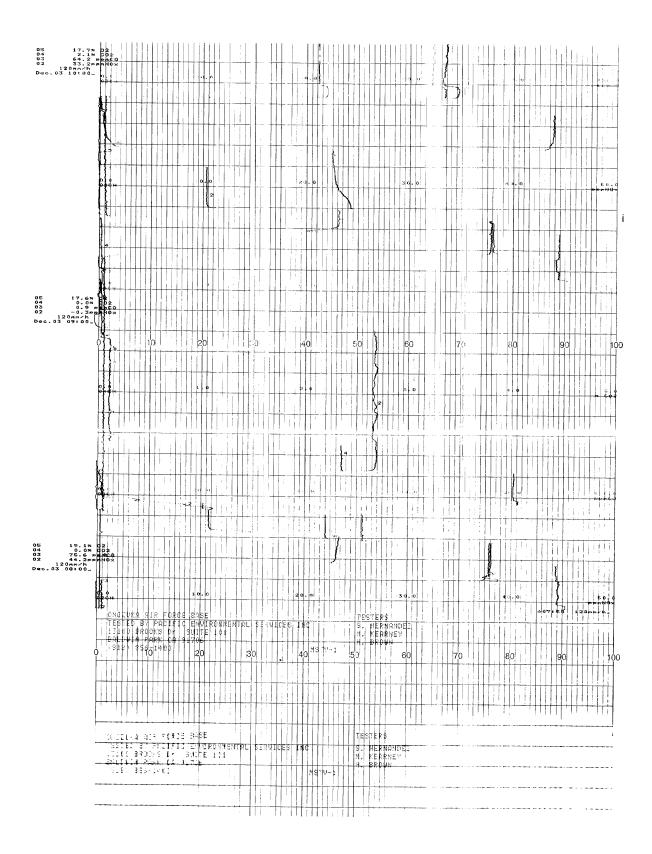
NOTE: NO SIGNIFICANT CATCHES

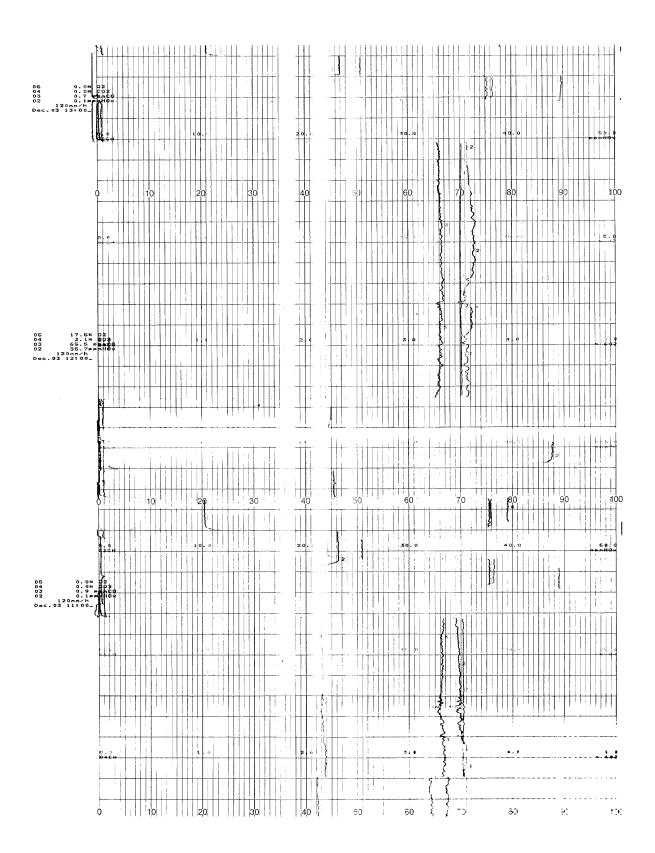
1.E. < 1 mg PM10 < 1 mg > PM10

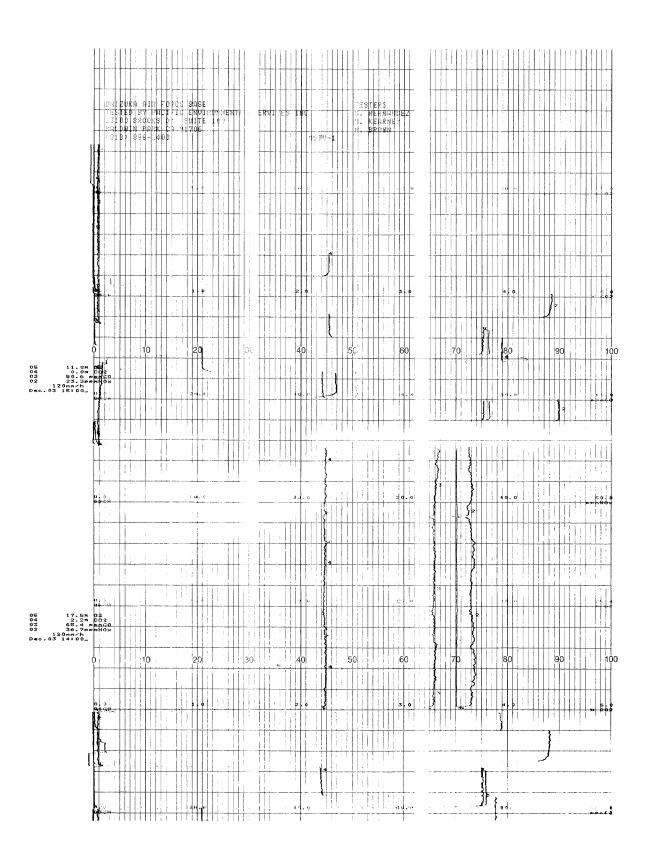
# FILTERS &

1	0.2084	0.2083	0.0001
2	0 2066	c. 2012	-0.0006
3	0.2056	0-2065	-0.0009









		Project No.	Page 1 of 8		
				<del></del>	EB
PACIFIC ENVIRO	NMENTA	L SERVICES, IN	ıc.	SUNNYYALE, C	Δ.
Prepared By	Date 12-ろ	Checked By Ы/Д	Da	e Sheet Title VAC & START 19.8"14	y - 19.8" e. GAD DIN 1 (CARB 20)

TIME	Nox	co	COL	02	NOTES,
	0.0	0,9	0.0	0.0	200 e 0.0 6 %09120
	44.2	75.6	41,0	19,0	H-SPAN 44.4
	23.1	51.0	2.3	11,0	M. SPAN 22-3
	10,7				L-SPAN 10.4
08:23	24.8				CONVERTER CHOR (NO.)
08:53	26.5	<del>.</del> .	 		6-10 CONVERTER CHECK
08:26		. <b>.</b>		17.6	OL TRAVORSE R 8 (4) SE
				17.6	2/ R7 (3)
_ <del>-</del> -		. — —		17.6	FC (2)
				17.6	R (2) R 5 (1) R 7 (4) NE R 3 (3)
				17.6	RY (4) NE
				17.6	3/10° R 3 (3)
				17.7	R 2 (2)
				17.6	/ R ( (1)
, <b></b>	43.8	75.5	2,2	19.0	Bias (High)
577427	eun 1				
09:56	34.7	لولا. ك	7.1	17.5	e R4 F/5 CO2= 15%
09:57	34.7	64.6	21	17.5	e 0.250 VOC
01:58	34.0	65.2	2-1	17.6	•
D9.59	<del>3</del> 3, 1	64.2	2.1	17.7	
€0100	33. 4	64.2	2.1	17-7	
10.01	33, 2	44.1	2.1	17.7	
10:02	<del>3</del> 3.3	64.2	2.1	17.7	
10:03	33,4	64.4	2.1	17.7	
10.64	33,5	64.4	2.1	17.7	
10:05	33,4	64.1	2.1	17.7	

					Project No. Page of 2 8				
				CI	Client ONIZUKA AFB				
PACIFIC ENVIRONMENTAL SERVICES, INC.					Sunguere, LA				
Р	repared By	Date	Checked By	Date					

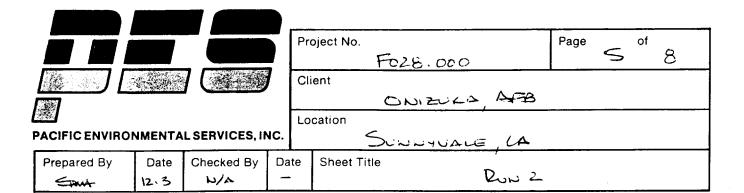
TIME	Nox	<u>co</u>	_CO2_	0_	NOTES:
10:06	33.5	64.4	科	13,7	
F0:01	33.5	64.3	2.1	15-5	
10108	33,5	64.3	211	127	
10109	33.5	64,3	2.1	12.7	
10:10	33,6	64.1	2.1	17,7	HADUSS to R 3 SW
10:11	33.6	64.2	2.1	17.7	
107.12	33.6	64.2	2.1	17.7	
10:13	· 33. 7	<b>64.3</b>	2.1	17.7	
10:14	35.7	64.1	2.1	17.7	
10:15	33.6	63.8	2.0	1ナナ	(20 niu)
10:16	33.5	63.9	2.1	17, 7	
10:17	33.7	64.1	2.1	13.7	SCTUAL POINT CHANGE
10:18	34.6	65.2	2.1	17.6	4 CHANGES OBSERVED ON
10:19	35.2	66.4	2.1	17.6	STEIP CUART.
10:20	35.3	46.7	2.1	17.6	
10:21	35.5	66.9	2.1	17,5	
10:22	<del>35</del> . 4	66.6	2.1	17.5	
10:23	35.5	64.5	2.1	17.5	
10:24	<del>3</del> 5.5	44.5	2.1	17.5	
10: 25	35.5	66,5	2.1	นเร	(30 MIN)
10:26	35.3	44.5	2-1	17.2	
10127	35.2	66.5	2.1	17.6	
10:28	55.1	Lele. Le	2.1	13.5	
10129	35.0	66.6	2.1	17.5	and the second s
10130	34,7	66.3	2.1	17.6	and the second s
10:31	<i>3</i> 4. 7	66.0	2.1	17.6	and the second of the second o
10:32	<b>3</b> 5.0	66.0	2.1	176	and the second s
10133	35.0	66.0	2.1	17.6	
10:34	34.6	65,6	2.1	17.6	
10.35	34.9	45.8	211	17.0	(40 Mm)

				roject No.	Page of 8
				CNIZUKA AFB	
				ocation	
PACIFIC ENVIRONMENTAL SERVICES, INC.			IC.	SUNDAULE, CA	
	Date	Checked By	Date	Sheet Title	
- Vi	2.3	N/A	-	Puni	·

TIME	Nov	<u>co</u>	<u> </u>	0_	Notes:
10:36	34.1	45.8	2.1	17.0	
10:31	349	45.9	2	17.6	
10:38	34. E	65.9	2.1	17.6	4 4000 40 Pt 3
10:39	34.8	65.8	2.1	17.6	
10:40	34.8	<b>66.</b> 0	2.1	17.6	
10:41	34.9	66.2	2.1	17.6	
10:42	34.8	662	2.1	17.6	
10:43	34.8	66.4	2.1	17,6	
<b>।</b> ८३५५	34.9	66.3	2.1	17.5	
10:45	34.8	46.3	2.1	17.6	(50 MIN)
10:46	34.8	<b>ل</b> ولو، ٤	2.1	17.5	
10:47	<b>3</b> ५. ት	66.2	2.1	34.5	
10:48	34,7	ک یایا	2.1	17.5	
10:49	34.8	46.2	2.1	17.6	
10:50	34,7	662	2.1	けん	
lo:51	34.7	46.0	2.1	17.6	,
10:52	34.7	أدلورا	21	17.6	$\begin{cases} \overline{X} & CO = \\ \overline{X} & CO_2 = \\ \overline{X} & O_2 = \\ \overline{X} & O_2 = \end{cases}$ $(CO MIN) \left\{ \overline{X} & NO_2 = \\ \overline{X} & O_3 = \\ \overline{X} & O_4 = \\ $
10:53	<b>うり</b> んし	١٠٠١ عاب	2.1	17.6	× 602=
10:54	34.6	44.4	2.1	17.5	\ \( \nabla \cdot \cdot \cdot \)
10:55	34.4	46.4	2.1	17.6	(LO MIN) { X NOX=
	C, 2	1.0	Ġ. ŭ	Ö. 0	300c
	44.4 7	5,4	4.0	19.0	H-SPAN
	23.1	D. 8	2.2	11.0	1492 - N
	10.2				4927
	43.7	15.5	2,2	19.0	BIAS

			Pr	Project No. Page of 8				
				Client ONIZUKA AFB				
PACIFIC ENVIRO	NMENTA	L SERVICES, II		SULLNYVALE, LA				
Prepared By	Date	Checked By	Date ~	Sheet Title	<u>-</u>			

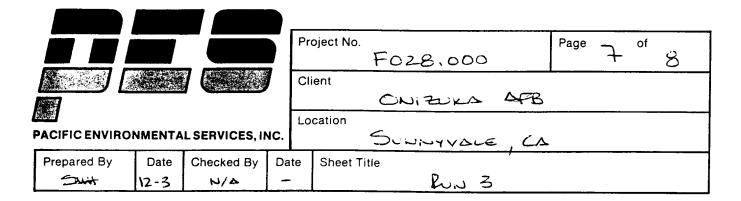
TIME	Nex	w	COL	<u> </u>	Notes
11:51	35,6	65.9	2,2	12.5	STAPT TEST PLY SE PORT
11:53	35.6	ليل. ٥	2.2	17.5	
11:31	35.6	45.8	2.1	17.5	
11:55	35.6	45.7	2.1	17.5	
11:56	35.6	65.8	2.1	17.5	
11:57	35.8	65.9	2.1	17.5	
11:58	<b>3</b> 5. 8	45.8	2.1	17.5	
11:59	<del>3</del> 5,7	45.5	2.1	17.5	
12:00	35.6	45.5	2-1	17.5	
12:01	35.6	43.7	2-1	17.5	(10 MIN)
12:02	35.6	46.0	2.1	17,5	
12:05	35.6	46.0	2.2	17,5	
12:04	35.5	66.0	2.1	17.5	
12:05	35·6	66.0	2.1	17.5	
12:06	35,7	66.0	2.1	17.5	
12:07	34.0	65.9	2.1	17.5	
12:08	36.1	66.0	2.2	17.5	
12:04	34.0	46.0	2.1	17.5	The state of the s
12:10	36.0	لول,ع	2.1	17.5	; ;
12:11	36.1	66.0	2. \	17.5	(20 MIN) [MOVE to 12 3] *
12:12	35.5	44.5	2.1	17.5	and the second of the second o
12:13	<i>35</i> .3	66.7	2.1	17.5	
12:14	35.5	46.5	2,1	17.5	
12:15	35.5	46.5	2.1	17.5	and the second s
12:16	35.3	46.5	2.1	17.5	and the second s
12:17	45.5	35.5	2.1	17.2	
12:18	35.9	لون, ٧	2.1	17.5	. <del>.</del>
12:19	<del>25</del> , 7	66.7	٤.١	17.5	
12:4	36.1	66,4	2.(	17.5	
12:21	36.2	نوله،۱	2.1	17.5	(30 MIN)



TIME	Nox	<u>co</u>	<u>cc</u> _	0-	NOTES.
12:22	36-3	66.2	2.1	17.5	
12:23	36.4	66-2	2.1	17.5	
12:24	36,4	46.1	2.1	17.5	
12:25	36.2	66.1	2.1	17.5	
12:26	36.2	46.4	2,1	17.5	
12:27	36.3	46.3	2.1	17.5	
12:28	36.4	66.2	2.1	17.4	
12:29	36.2	ا ، <i>حاجا</i>	2.1	17.5	
12130	36.2	66.2	2.1	17.5	
12131	36.1	46.2	2.1	17,5	(40 MIN) (SWITZERED & RZ)
12:32	36.1	66.2	2.1	17.5	
121 53	34.0	66.1	2.1	17,5	
12:34	36.0	46.0	2.1	17.5	
12135	36.1	45.9	<b>2</b> , ι	17.5	
12:36	3φ.(	44.0	2.1	12.5	
12:37	36.1	65.7	2.1.	17.5	
12/38	36.1	45,8	2.1	17.5	
12:39	36.0	65.B	2.1	17.5	
12:40	36.0	45.9	2.1	17.5	
12:41	35.9	45, 8	2-1	17.5	(D) MID)
12:42	35.8	65.7	2.1	17.5	
12: 43	<i>3</i> 5.8	65.9	2.1	17.5	,
12:44	35.8	45.8	2.1	17.5	
12: 45	35,7	65.8	2.1	17.5	
12:46	35.7	45.8	2.1	17-5	
12: 47	35.6	458	2.1	17.5	
12148	35.6	65.5	_	17.5	
12:49		45.9		17.5	
12:50		65,8		13,5	
12:51	35. L	<b>6</b> 5,8	2.1	17.2	(60 MIN)

			Pr	Project No. Page of 6 8				
			CI	Client ONIZUKA APB				
<b>1</b>			Lo	Location				
PACIFIC ENVIRO	NMENTA	L SERVICES, IN	IC.	SUNDAVALE LA				
Prepared By	Date	Checked By	Date	Sheet Title				
Prepared By	12-3	N/A	-	Eun 3				

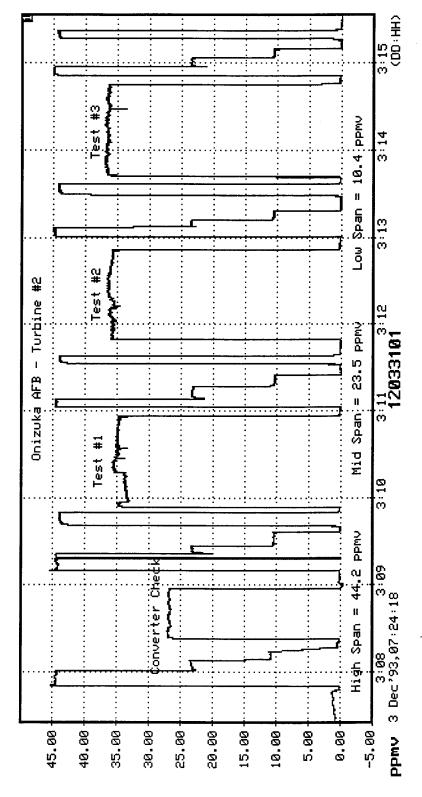
TIME	Nox	<u>ao</u>	_co	O_	Notes:	
_ ~	0.1	0,7	-0.1	0.0	Zero	+0.01
	44,6	75.2	3,8	19.0	H-SPAN	3.9%
مطاهم حدود	23.3	30.5	2. \	llio	M-SPAN	
	10.3				L-SPAN	
	44.0	75.0		18.9	Bias	
		Manda in the stript in the street was	Andrew Control of the			
			E METHOD I I I I I I I I I I I I I I I I I I I			
13:46	36.4	65.5	2,2	12,4	STAR	- AT R 2 SE
13:47	36.5	45.5	2,2	17.5	1	
13:48	3666	65.5	2,2	17.5		
13:49	36.6	65.6	2.2	17.5		
131:50	36.7	65.6	2.2	17.5		
13:51	36,7	65. F	22	\$ 17.5		omenic color accommendado. En como acomo escaba con
13:52	36.6	45.6	2. 上	17.5		
13:53	36.6	45.7	2,2	12.५		AND AND AND AND AND AND AND AND AND AND
13:54	34.6	45.6	2.2	12,4		The state of the s
13:55	36.7	65.6	2.2	17.4	(10 MIN)	
13:56	36.8	65.8	2,2	14.4		and the contract of the contra
13:57	36.6	65.5	2,2	17.4		
13:58	36.6	45.6	2.2	12.4	3	
13.59	36.6	45,5	2.2	14.5	And the second s	enne mario alla con esta esta esta esta esta esta esta esta
14:00	34.6	65.4	2,2	17-5		and the second s
14:01	B6.5	45.3	2.2	17.5		en en en en en en en en en en en en en e
14:02	36.5	65.5	2.2	17.5	: :	••••
14:03	36.6	65,6	2.2	17,5	and the second of the second o	And the second s
14:04	36.6	65.6	2.2	12.5		
14:05	36.5	45.5	2, 2	1チ、ら	(20 MIN)	MULD TO P3 SE

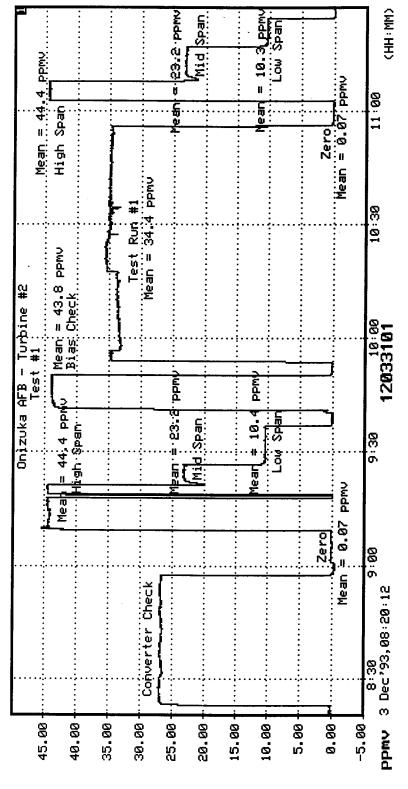


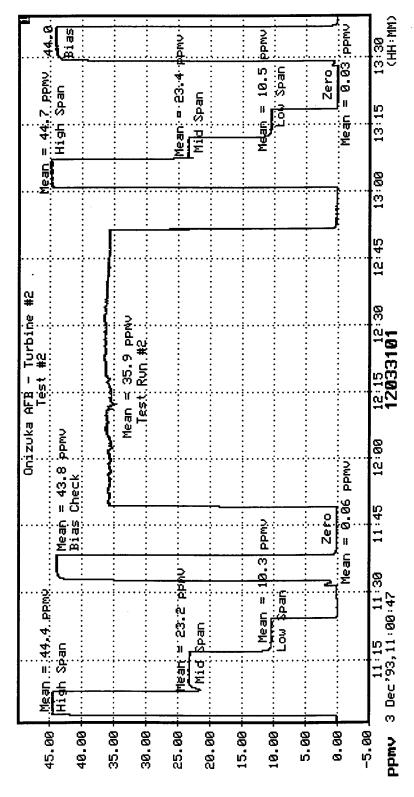
TIME	NO+	<u>co</u>	COL	0_	HOTES.
14:06	34.3	65.5	22	17,5	
14:07	36.2	65.1	2.2	17.5	
14108	36.3	45.3	2,2	17.5	
141.09	36.3	45,5	2,2	17,5	
Millo	36.3	45.6	2,2	17.5	
14:11	<i>3</i> 4.3	65.4	2.2	17.5	
14:12	36.3	65.5	2.2	14.5	
14:13	36.4	45.5	2.2	17.5	
14:14	36.5	45.4	2.2	17.5	
14:15	36.4	45.5	2.2	17.5	(30 MID)
14:16	36.3	454	2.2	17.5	
14:17	36.5	45.5	2.2	17.5	
14:18	36.7	45.6	2.2	17.5	
14:19	36-7	45.7	2.2	17.5	
14:25	36.5	65.8	2.2	17,5	
14:21	36.5	45.6	2.2	17,5	
14:22	36.7	65.5	2.2	17.5	
14.23	36.6	45.5	2.2	17,5	
14:24	36.6	65.5	2.2	17.5	
14.25	36.4	45.5	2.2	17.5	(40 MIN) MINED to PLY SE
14:26	36.4	45.5	2.2	17.5	-
14:27	36.3	65.5	2,2	17.5	
14:28	36.2	65.5	2.2	17.5	•
14:29	36.2	45.5	2,2	17.5	
14:30	36.0	<b>43.7</b>	2.1	17.6	PROBE HOURD FOR ANIO
14131	36.2	65.6	2.2	17.5	
14:32	36.0	65.6	2.2	17.5	
14:33	36.3	<b>65</b> .8	2,2	17.5	
14:37		65.U		17,5	
14:35	36.5	<i>45</i> ,8	2,2	17,5	(50 MIN)

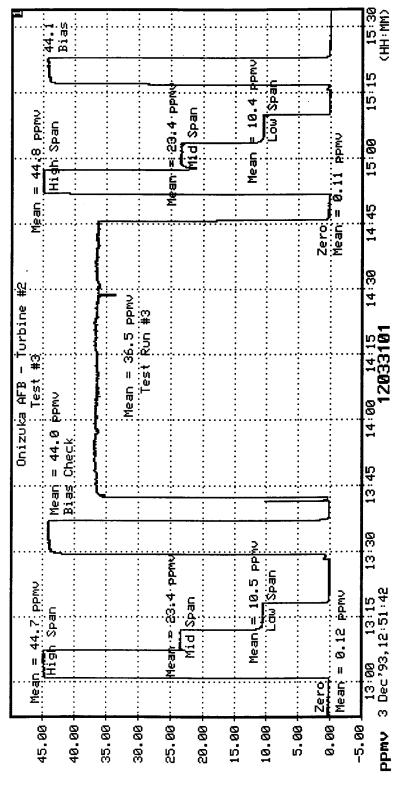
			Pı	roject No.	28,000	Page o	f 8
				lient	ONIZUKA	AFB	
PACIFIC ENVIRO	ONMENTA	AL SERVICES, II		ocation	SUNNYYAUE	, LA	
Prepared By	Date	Checked By	Date	Sheet Title		N 3	

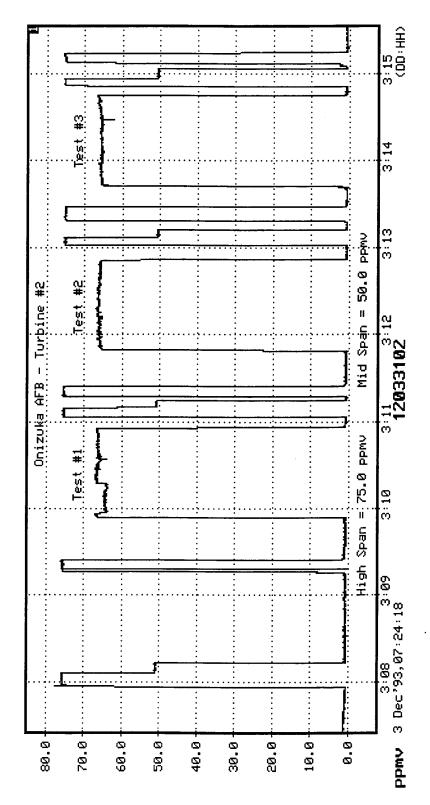
TIME	Non	Co	CO2	02	NOTES:
14:36	36.5	65.9	2.2	11,5	
(4:37	36.5	66.0	2,2	17.5	
14138	36.4	65.8	2,2	17.5	
14139	364	66.1	2.2	17.5	
14:40	36.4	66.2	2.2	17.4	
14:41	36.2	46.3	2.2	17.5	
14:42	36.2	66.2	2.2	17.5	
14:43	36.3	66.0	2.2	17.5	
14:44	36.3	66.2	2.2	17.5	
14:45	36.2	46.4	2.2	17,5	(60 MIN)
				·	
	C . 6	0,7	0.0	6.0	26%
	44.8	75.1	3.9	19.1	H-SPAN
	23.3	50.5	2.2	11.0	H-SPAN
<del>-</del> -	10.4		-,-		L-SPAN
	43.7	75.0	2.2	19,0	Bias.
15;-	44.1			TO THE SECOND AND THE SECOND ASSESSMENT	END OF TEST.

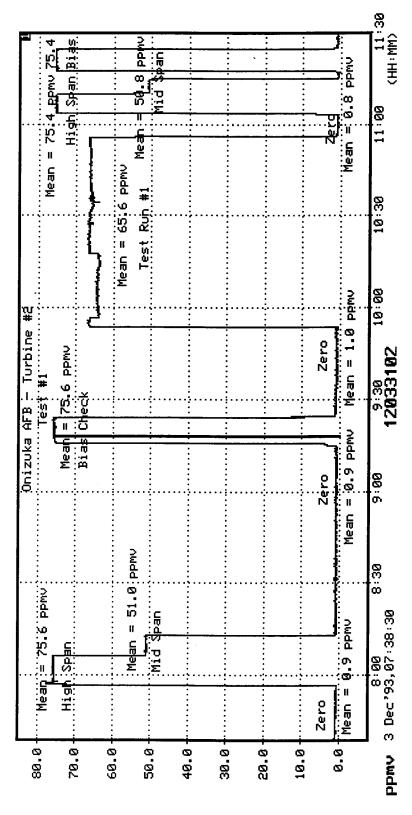


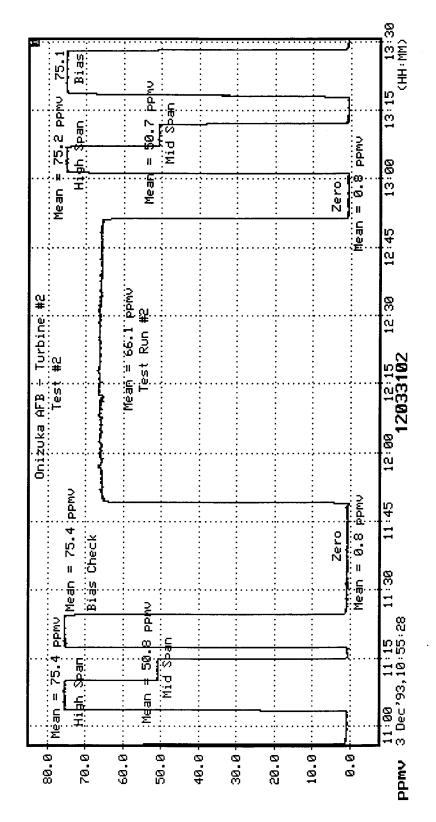


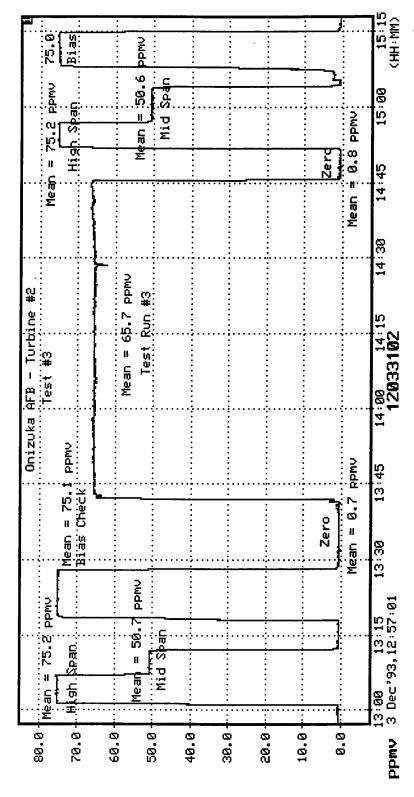


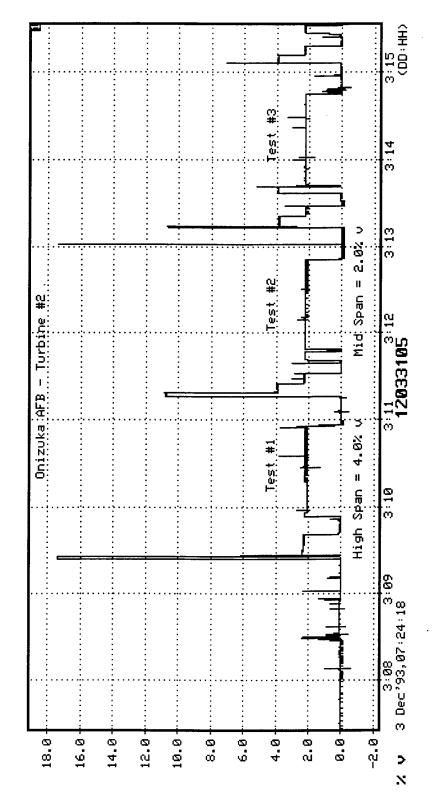


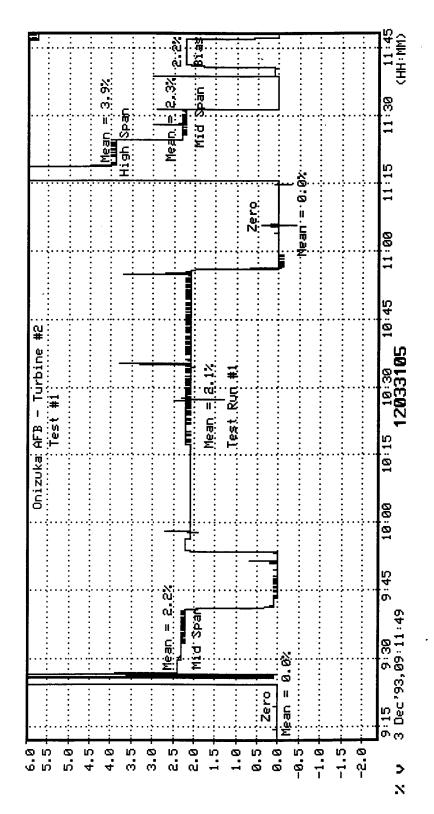


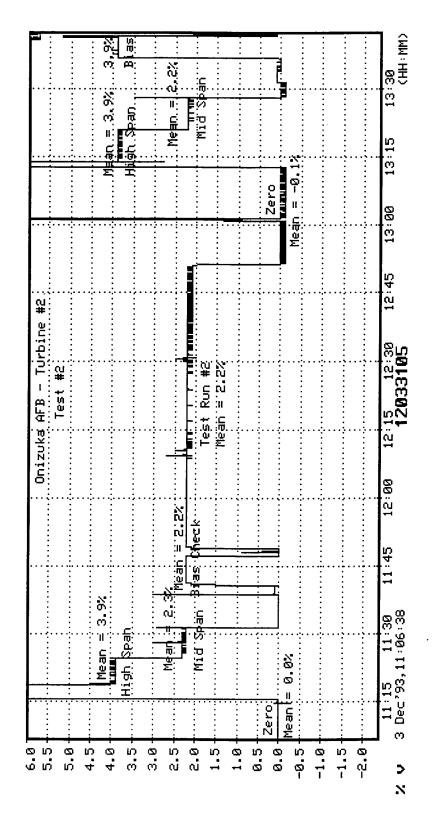


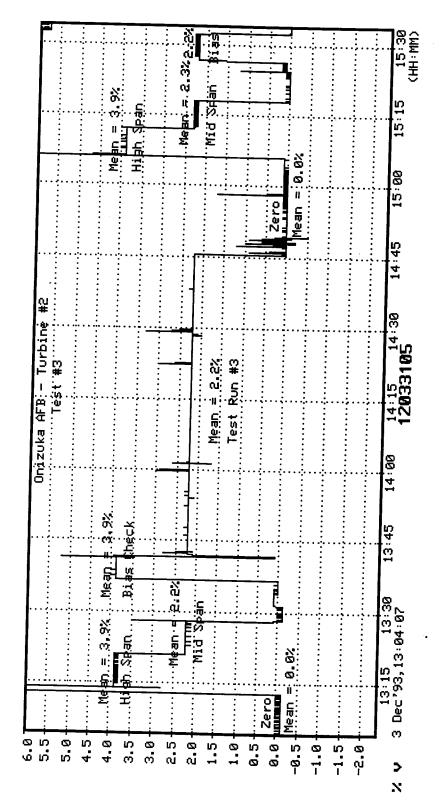


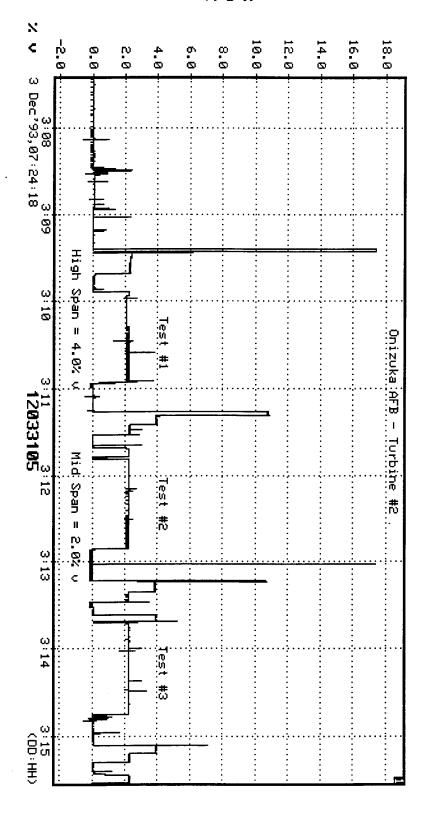


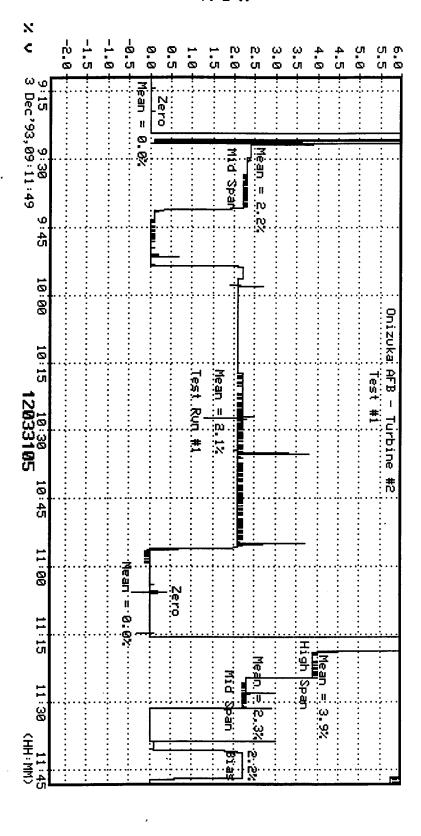


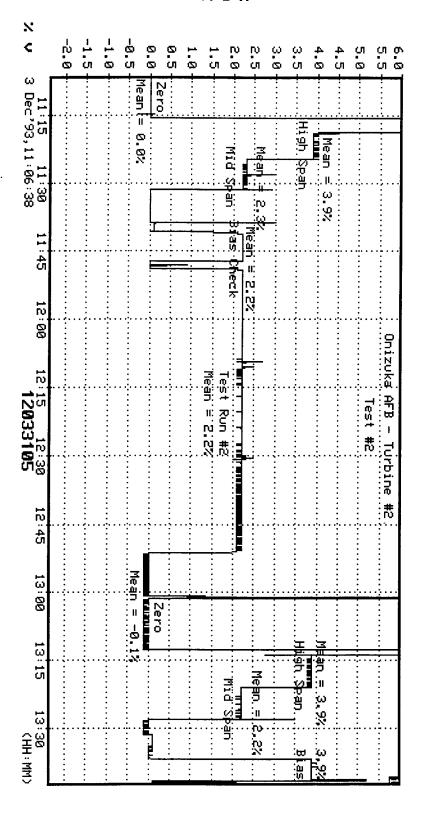


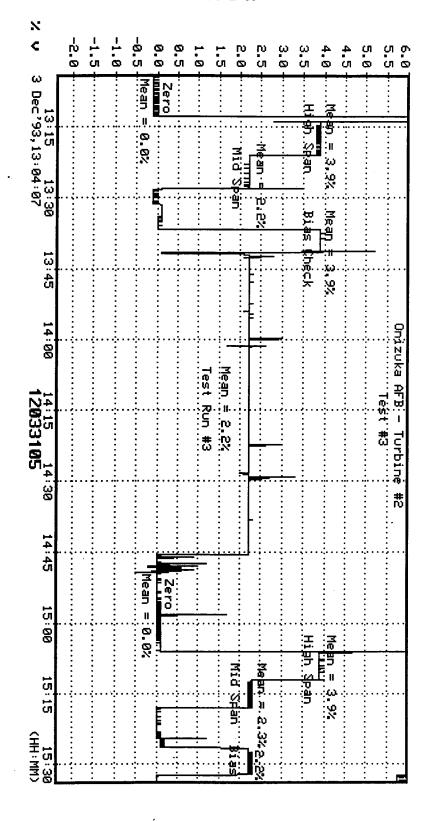


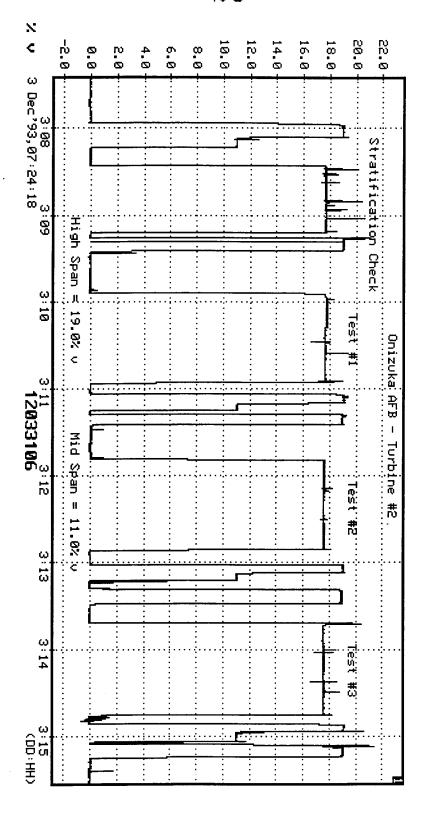


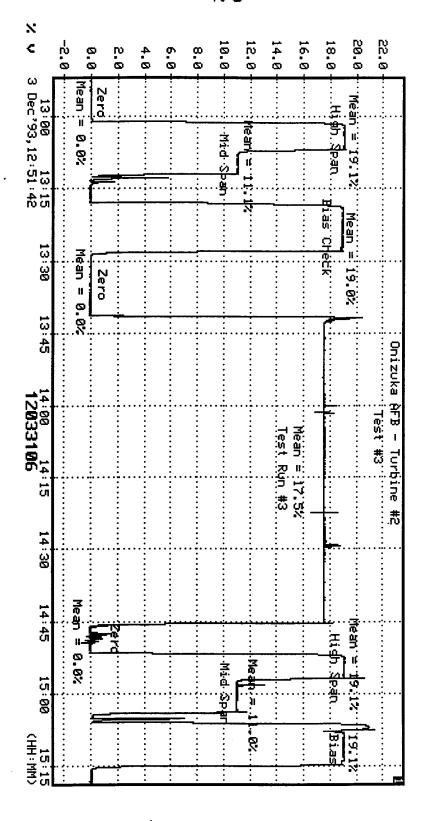














Project No. FOZ8

Client

ONIZUKA AFB

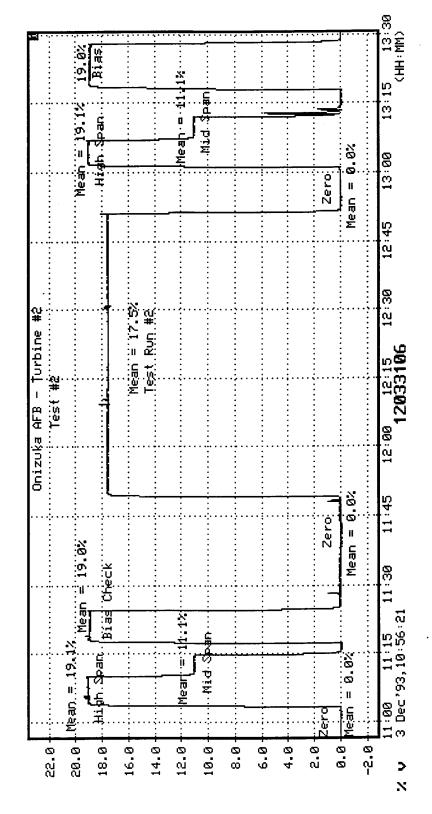
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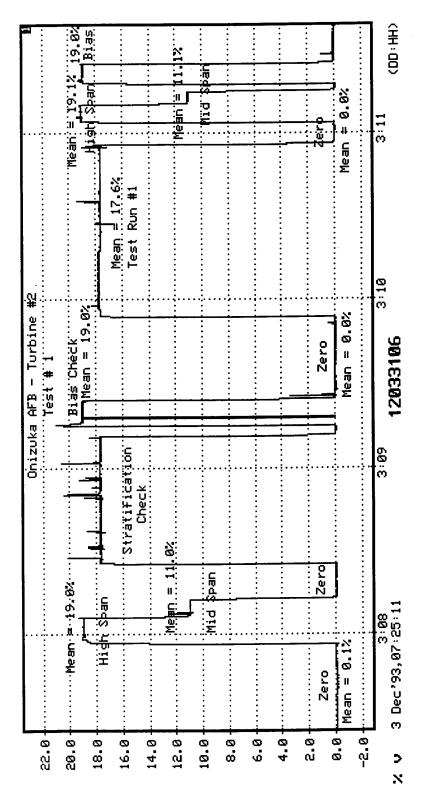
TURBINE #2 EXHAUST

PACIFIC ENVIRONMENTAL SERVICES, INC.

repared By Date Checked By Date Sheet Title CONTINUOUS MONITORING QA/QC

,		PRE-	TEST	POST-T	EST	
NOx	SPAN VAWE PPMV	response PPMV	CAL ERROR %F.S.	RESPONSE PPMY	CAL ERROR % F. S.	DRIFT % F.S.
	and the state of t					COMMENT OF THE COMMEN
TEST *1	The second secon				10.11	
ZERO	0.00	0.07	0,14	0.07	0,14	0,00
LOW SPAN	10.4	10.4	0.00	10.3	-0.20	-0.20
MID SPAN	23,5	23.2	-0.60	23,2	-0,60	0.00
HIGH SPAN	44.2	44.4	0.40	44.4	0,40	0,00
BIAS	44.2	43.8	-0,80	-		
		3	1			
TEST #2						
ZERO	0.00	0.06	0.12	0.03	0.06	-0,06
LOW SPAN	10,4	10.3	-0.20	10.5	0.20	0,40
MID SPAN	23.5	23.2	-0.60	23.4	-0.20	0.40
HIGH SPAN	44.2	44.4	0.40	44.7	1.00	0.60
BIAS	44.2	43.8	-0,80	-	-	-
	:	The second secon	1			
TEST #3	***************************************					
ZERO	0.00	0.12	0,24	0.11	0, 22	-0.02
LOW SPAN	10.4	10.5	0.20	10.4	0.00	-0.20
MID SPAN	23.5	23.4	- 0.20	23.4	-0.20	0.00
HIGH SPAN	44.2	44.7	1.00	44.8	1,20	0.20
BIAS	44.2	44.0	-0,40	44.1	-0.20	. 0,20
	, , .	•	· · · · · · · · · · · · · · · · · · ·			





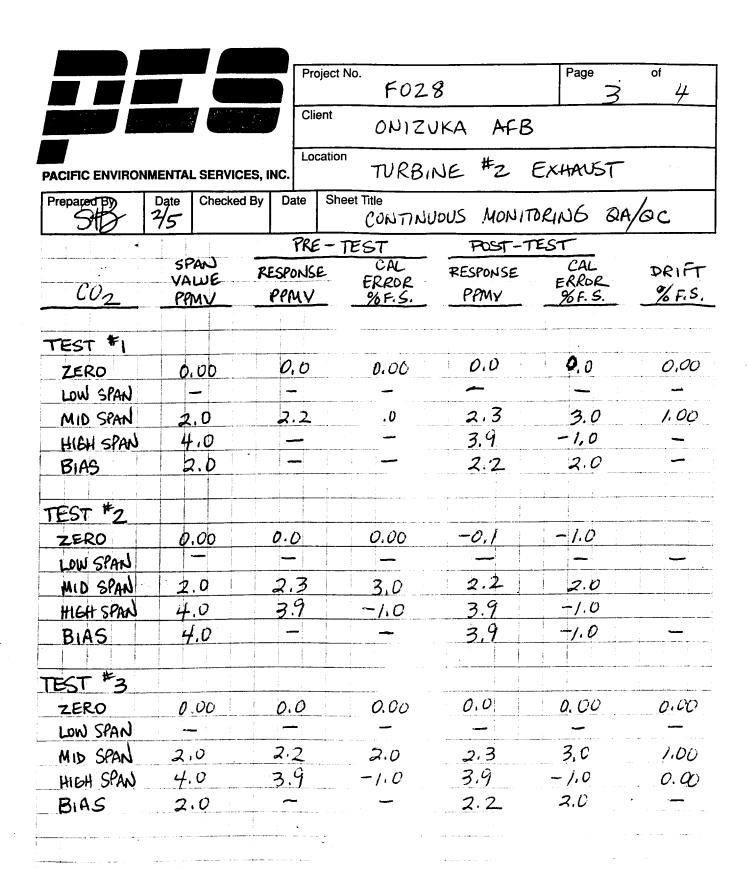


Project No	F028		Page	2_	of
Client	ONIZUKA	AFB			
Location	TIPRINE	#2. Ex	HAUS	7	

PACIFIC ENVIRONMENTAL SERVICES, INC.

Sheet Title CONTINUOUS MONITORING DA/QC

Prepared By	Date Check	ed By Date She	eet Title CONTNU	IOUS MONITI	DRING BI	4/oc
		PRE-	TEST	POST-T	EST	
CO	SPAN VALUE PPMV	response PPMV	CAL ERROR %F.S.	RESPONSE PPMY	CAL ERROR % F. S.	DRIFT %F.S.
TEST *1	The second secon	Account of the second property of the second				
ZERO	0.00	0.9	0.90	0.8	0.80	-0.10
LOW SPAN		-	-		CONTRACTOR CONTRACTOR STORE TO THE STORE S	
MID SPAN	50.0	51.0	1.00	50.8	0.80	-0.20
HIGH SPAN	75.0	75.6	0.60	75.4	0.40	-0,20
BIAS	75.0	75.6	0.60			
The second secon			BELLEVINOR OF THE STATE OF THE			
TEST #2						
ZERO	0.00	0.8	0.80	0.8	0.80	0.00
LOW SPAN						
MID SPAN	50.0	50.8	0.80	50.7	0.70	-0.10
HIGH SPAN	75.0	75.4	0.40	75.2	0.20	-0.20
BIAS	75,0	75.4	0.40	-		
TEST #3				1		
ZERO	0.00	0,7	0.70	0.8	0.80	0.10
LOW SPAN	<b>-</b>					
MID SPAN	50.0	50.7	0.70	50,6	0,60	- 0.10
HIGH SPAN	75.0	75.2	0.20	75.2	0.20	0.00
BIAS	75.0	75.1	0.10	75.0	0.00	





Date 2/5

Project No	F028		Page	4	of #
Client	ONIZUKA	AFE	3	-	
Location	7100 16	‡ <sub>2</sub>	EVILANT		

Checked By Date Sheet Title CONTINUOUS MONITORING DA/OC

•		PRE-	TEST	POST-T	EST	
02	SPAN VALUE PPMV	response PPMV	CAL ERROR %F.S.	RESPONSE PPMY	CAL ERROR %F.S.	DRIFT %F.S.
		1				1 1
TEST *1						
ZERO	0.00	0.1	0.40	0,0	0.00	-0.40
LOW SPAN						
MID SPAN	11.0	11.0	0.00	11.7	0,40	0.40
HIGH SPAN	19.0	19.0	0,00	19.1	0.40	0,40
BIAS	19.0	19.0	0.00	_	-	-
TEST #2						
ZERO	0.00	0.0	0.00	0.0	0.00	0,00
LOW SPAN	-					
MID SPAN	11.0	11.1	0.40	11.1	0.40	0.00
HIGH SPAN	19.0	19.1	0.40	19.1	0.40	0,00
BIAS	19.0	19.0	0,00	-		_
1				1 - 1		
TEST #3						
ZERO	0.00	0.0	0.00	0.0	0.00	0.00
LOW SPAN				_		
MID SPAN	11.0	11.1	0.40	11.0	0.00	-0.40
HIGH SPAN	19.0	19.1	0.40	19.1	0.40	0.00
BIAS	19.0	19.0	0.00	19.1	0,40	
	•		· · · · · · · · · · · · · · · · · · ·			1 1111

750 SPTS/DE FORM 34, JAN 32 SEPLACES 10048958QUDE FORM 34, OCT 89, WHICH WILL BE USED PACKAGE NUMBER TOTAL TO DATE 2300 1800 **AS** 25 15 1400 410 130 2200 TOTAL TODAY PREVIOUS TOTAL 8 1900 8 1,700 14 667 1300 % 1100 4/60 000 100 26,077,000 26,050,000 026,250,000 1200 40 700 9 8 ENGINE OPERATING HOURS 0200 H/C 180 0100 11/2 Š ENGINE SERIAL NUMBER 727 ¥ DAILY POWER PLANTOPERATING LOG - (TURBINE - BOILER) ŝ Z KVAR 124774 3/ca dan 7 4 -₹ # 2 74 AMENIALIS T-7 | T-6 | PCD 10.00 OI. A000ED SOF Sec 0654 630 8% 240 820 64 87 088 089 É 140 820 69 650/8/0 69 69018089 630 870 45 820 890 0,0 8 80 68 8/0/8 20 SWINGS ي ý DAYS SQIN 3 3 2 75 26 77 ę Ð 140 1 140 727 2 801 /3/ 721 ē 0 75 Ý ۲, 2 2,3 TENE 83 58 10  $\overline{\varsigma}$ 4 1-12 ū 3 BOILER 5 4 PACKAGE NUMBER TOTAL TO DATE PREVIOUS TOTAL 2300 2200 2100 TOTAL TODAY 200 **8**00 1700 100 100 M. E 12 0081 8 1500 (2) 20 8 8 8 **3** 8 8 88 0700 800 9500 8 0200 ENGINE OPERATING HOURS 8 8 No. 3MI ENGINE MFR GENERATOR MFR ENGINE SERIAL NUMBER 520 570 09 068 520 5% 570 50 ₹ 0450 0 60 20 ç KVAR CO 13612 26 -¥P 57471 = 1 2 TURBINE AND GENFRATOR

AMPS AMPS VOLTS T-7 T-6 PCD ¥. ELECTRIC MACHINERY SWINGS DAYS SQIM 20646 OIL ADDED ž 700 960 200 7 8 940 920 γ 72 22 23 23 Z . د:ę MODEL 5 5ء BEMAC II 0 TANET ß Ç 2/ PACKAGE NUMBER 2300 1300 470 TOTAL TO DATE 33 2100 TOTAL TODAY PREVIOUS TOTAL 8 0700 480 8 1700 460 2001 **1**500 8 1000 /F/ JS ğ 1 200 8 1000 460 300 VOLTAGE 4160
ENGINE SERIAL NUMBER 8 0800 0500 1756 2nm | 0000 8 ENGINE OPERATING HOURS 8 0500 Ario 250 3MIL 8 Ň E 940 920 470 20 22,000 40 300 ₹ 30 8 300 250 300 000 KVAR 546611 der 6 % -A ź TURBINE AND GENERATOR × 12 00 +240 63 KW 750 MBS DAYS OIL AUDED 20636 280 087 120 800 400 130 370 080 087 20/07/38 マララ ナベ 8 000 880 870 80 CX 2010 8 20 ğ 36 36 75 ş 3 g 5 X Drc •ნ 3 c5512712 53 12.715 53 7.5 15 31/56/1 /25 25 34 12 -115 E AFE 35 ε اد 12.5 12.5 12.5  $\tilde{\mathcal{L}}$ 12 2 3 R 16

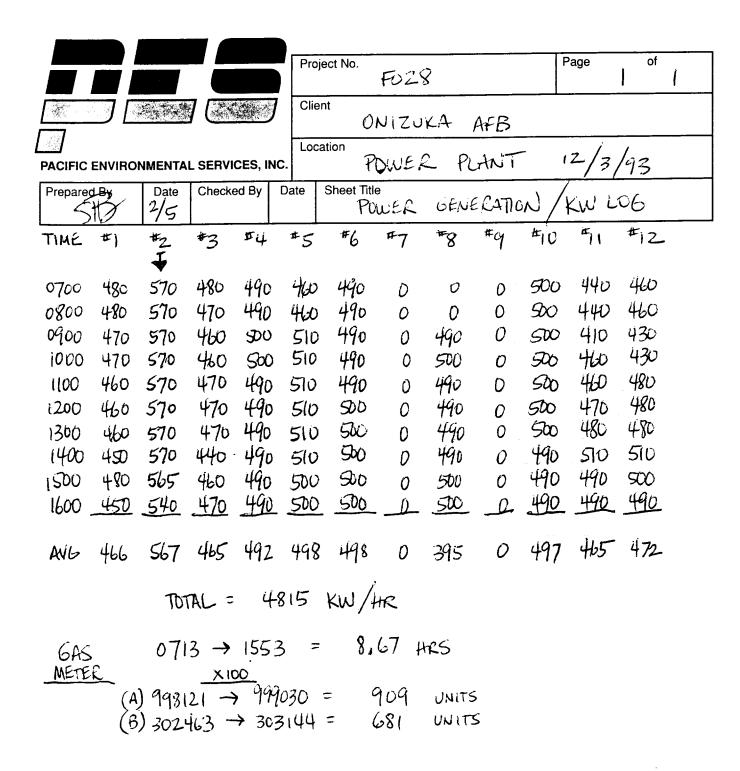
150 SPTS/DE FORM 34, JAN 92 REPLACES 1004SPSSQ/DE FORM 34. ACKAGE NUMBER TOTAL TO DATE TOTAL TODAY 2200 3100 PREVIOUS TOTAL 8 88 1600 490 20 8 176 Challed 24 1300 490 2,20 1200 196 220 1100 490 200 9.11-5.7 1000 الحدد أسم ا 2000 ENGINE OPERA 0/1/005/0010 ENGINE SERIAL NUMBER POD FERES 35 (25) 500 150 24 1 205 28 150 07112 ž DAILY POWER PLANT OPERATING LOG - (TURBINE - BOILER) KVAR 125000 ÿ 73 170 2 - 3 マ URBINE AND GENERATOR DAYS SGIN 유 VOLTS 1-7 | 1-6 PCD 7113617 2012/s 20 000 21 100 00 16 598 069 080 1850 122 152 130 15820 650 046 75 50 173 670 830 77 67 179 630 830 75 38 357 120 32 00 16 098 7 2 2 13 DAYS 77 57 126 SQIW 5 9 ナンサン 52 53  $\ddot{c}'$ 57 179 57 3 ę 02 OPERATORS 5 6 7 4 65 L \_ τ .: τ ٤ 164 S ڏ ট ンニュ 8 2 6 PACKAGE NUMBER 2100 TOTAL TO DATE TOTAL TODAY PREVIOUS TOTAL 2300 2200 2000 88 700 8 8 8 8 <del>3</del>0 1200 <del>1</del>8 8 8 0500 GENERATOR MFR
ENGINE SERIAL NUMBER 880 0700 386 0771 como 0200 0100 2911 0000 Š TIME 3 25 00 00 1150 910 110 ¥ 282 200 200 120 Ŗ, 130 130 KVAR -5 56 541 24 TURBINE AND CINITATOR # Ž  $\mathcal{C}_{\lambda}$ ELECTRIC MACHINI RY 8 SWINGS DAYS SQIM 읃 2(61) 13. 330 M 5 5 8 901830 171 3 200 7, 17.5 7 2 7/ 7 g 154 15 ٧, ę 56 176 MODEL 15 77 5 5ء BEMAC II TAN MARTIN Š 7 PACKAGE NUMBER TOTAL TO DATE 2300 200 1900 PREVIOUS TOTAL 2100 TOTAL TODAY 88 100 July 100 8 JOC 3000 VOLTAGE 4160
ENGINE SERIAL NUMBER 1/ 02/ 0001 8 30 1200 ī 8 0700 8 8 0500 5/0 700 8 020 ENGINE OPERATING HOURS 8 MON 3MIT 190 3 80 \$90 500 200 3, 500 000 490 Obh 50 ¥ Elen slenn 8 B 180 KVAR 700 190 20 35 139438 Z 20/4 AMPS AMPS AMPS # 0 TURBINE AND GENERATOR 3 KW 750 64522 SGIW SWINGS DAYS OIL ADDED 70 g 130 CSO 00 800 089 62 Cic 7 670 870 670 170 13 (11) 306 July 11; 60 300 80 890 37 Š 6. 1 ۲ 7 20 20 >. 2 2) > ğ **'** 65 . 3 Ē 111 : Š 16. Ę • 1 62 3 7 62 6. 3 ٤. ž. 13.00 7 2 7 .-3 17 0 74 12 5 13 1 2 7 ×

750 SPTS/DE FORM 34, JANEZ: EPLACES 1004SPSSQIDE FORM 34, OCT 89, WHICH WILL BE USED TOTAL TO DATE TOTAL TODAY PREVIOUS TOTAL 2200 200 900 8 10 1 8 8 1200 1100 8 0700 8 0400 300 0200 Š 3ML ENGINE SERIAL NUMBER × DAILY POWER PLANTSPERATING LOG - (TURBINE - BOILER) KVAR 12017 Ħ TURBINE USD GENERATOR SMAGS SAC **5**5/ 0612 SWINGS DAYS SGIM ę Ć 6 5 <u>ر</u> د ا hur ? 1 უნ Maria Maria Maria 3 BOILER REMARKS TOTAL TO DATE TOTAL TODAY 2300 2100 2000 88 1100 490 PREVIOUS TOTAL 2200 8 1800 1500 (7) 8 **1**00 8 <del>1</del>28 900 38 0700 GENERATOR MFR
ENGINE SERIAL NUMBER 888 800 0500 040 0300 ENGINE OPER 0200 Š ĦME 5 490 190 470 ,80 190 ¥ 7 180 on line & 850 180 KVAR AMPS AMPS 138395 Ħ 20 20 SOLAR
ELECTRIC MACHINERY TURBINE AND GENERATOR

MPS AMPS VOLTS 1.7 16 PCD 22 SWINGS DAYS SQIM JO 970 190 390 131 37 4 3 ١ 4 ę MODEL 15. 12 44 12 5 2.5 57 5ء BEMAC II T-1021S-21 Š BOILER 6 26 PACKAGE NUMBER REMARKS 2200 TOTAL TO DATE PREVIOUS TOTAL 2100 2000 TOTAL TODAY 8 8 1700 8 ğ 9 0800 8 00 ENGINE OPERATING HOURS 8 300 <del>1</del>28 100 0800 8 VOLTAGE 4160
ENGINE SERIAL NUMBER 0000 8 0300 0200 0100 MDN JWE ş ZVAR 144810 # TURBINE AND GENERATOR 0 KW 750 20658 SWINGS DAYS SQIM OIL ADDED ğ ē Ģ W DEC შ TEMP IN 8 BOILER

PACKAGE NUMBER #/C PACKAGE NUMBER #/C PACKAGE NUMBER #/C PACKAGE NUMBER #/C	Alban organi + 746	SWINGS	E SUNGE DATE DE LET	51337 mass Mins 1104	OPERATORS				2000	0000		77 14 7 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2	X 17 0 14 10 17 14	120 180 120 120 120 120 120 120 120 120 120 12	200 200 100 100 100 100 100 100 100 100	200 800 72 57 139 Ø 65 12 14	200 May 700 840 12 57 138 0 63 13 16	520 200 100 100 100 100 100 100 100 100 1	500 100 120 122 13 72 76 870 13 56 137 P 56 13 16	200 870 74 56 176 9 57 13 16	200 July 200	010 1/2 125 12 12 1/2 1/2 1/2 1/2 1/2 1/2 1/2 1/2 1	1/20 12:21 5 13:20 50 13:20 13:21/2	500 1170	510 170	170 70 71 70 700 800 700 800 150 155 8 54 12716	31 561 73 E + 21 65 HG 098 006	TIME KW KYAR AMPS AMPS AMPS AMPS COLTS T.7 T.6 PCD LOP LOT LO INLET PS P		ENGINE SERIAL NUMBER	DAILY POWER PLANT OPERATING LOG - (TURBINE - BOILER)
PACKAGE NUMBER # //	unt bourg down or	TOTAL TO DATE SWINGS	TOTAL TODAY DAYS	157157 1	ENGINE OPERATING HOURS OIL ADDED	6		8		8	17 (7) (0) 5/0 fe 70 70 10 10 1 1 1.	2 6	150 000 00 00 00 00 00 00 00 00 00 00 00	510 210 210 210 210 210 21 110 4 21 1: 1	70 54 140 4 70 19	420 180	51 67 6 AM 65 AL ALB AR	170 62 100 100 100 100 100 100 100 100 100 10	70 120 64 63 64 163 16 16 17 18 18 18 18 18 18 18 18 18 18 18 18 18	21 54 136 4 59 13		11 17 14 26 15 16 CON 19 19 19 19 19 19 19 19 19 19 19 19 19	100 0 1 7 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	C C C C C C C C C C C C C C C C C C C	100 100 100 100 100 100 100 100 100 100	10 100 10 10 10 10 10 10 10 10 10 10 10	TO COT COT COT TO TAKE PS	TURBINE AND GENERALLIN	ı	ENGINE SERIAL NUMBER  ELECTRIC MACHINEN	SOLAR T-10
PACKAGE NUMBER # $/2$	REMARKS  Then ober 1 these Til	TOTAL TO DATE SWINGS	DAYS	PREVIOUS TOTAL 119276 MIDS 8CAPLE # 2050	GINE OPERATING HOURS	2200	2200	300	1900		19 0 (2) (2) 24 06 20 14 00 10 10 10 00 00 00 00 00 00 00 00 00	12 1 1 02 16 8 DL	72 C C C C C	200 510 210	130 HAD 200 CH 1 20 1 2 1 2 1 2 1 2 1 2 1 2 1 2 1 2 1	1200 1/80 200	430 130 71 57 132 1 480 200	1000 1100 110 10 10 10 10 10 10 10 10 10	22 15 1 05 25 000 25	0700		0000 450 150 6 60 60 73 75 130 0 50	140 100 100 100 100 100 100 100 100 100	x 3 30	15 0000 44 160 62 62 65 850 850	15 mon 1140 160	P TIME KW KVAR AMPS AMPS VOLTS 1-7 1-8 PCD LOP LOT LO INLET	TURBI	48ER 2069/	VOLTAGE 4160 KW 750	KW 750

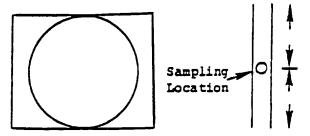
1500 17



4815 KWH  $\times$  8.67 HRS = 41746 KWH 41746 KWH / 909 UNITS = 0.0218 WITS/KWH 41746 KWH / 681 UNITS = 0.0163 UNITS/KWH

## PRELIMINARY VELOCITY TRAVERSE

Plant:	ONIZUKA AFB
Date:	1-12-94
Location	
Stack I.	
	ic Pressure, in. Hg: 30,22
Stack Ga	uge Pressure, in. H2O:O./
Operator	
Pitot Tu	be I.D. Number: ST4
Temperat	ure Readout I.D.: FISHER/TC-12
Pitot Tu	be Leak Check: OK



Schematic of Traverse Point Layout

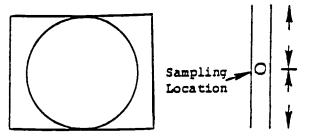
	: CA 59			
	Velocity	Stack	Cyclonic	
Point	Head $(\Delta p_g)$	Temp.	Flow Check	
Number	in. H <sub>2</sub> O	(Tq), °F	° from Null	
<u> </u>	1.40	450	1.6	
2	1.25	451	6.0	
3	1.60	438	0,8)	
4	1.25	434	22.3	
		443		
	ľ			
Average				

Traverse	Velocity	Stack	Cyclonic
Point	Head (Apg)	Temp.	Flow Check
Number	in. H <sub>2</sub> O	(T <sub>q</sub> ), °F	• from Null
RI	1.40	444	
2	1.45	450	
3	1.55	449	
4	1.25	444	
-			
		447	
		•	
	V		
Average	1,18	445	
		<u>-</u>	

- PACIFIC ENVIRONMENTAL SERVICES. INC. -

## PRELIMINARY VELOCITY TRAVERSE

Plant:	ONIZUKA AF	B
Date:	1-12-94	
Location:	TURBINE #2	STACK
Stack I.D.:		
Barometric	Pressure, in. Hg:	30,22
Stack Gauge	Pressure, in. H	
Operators:	<u>B</u>	ROUN
	I.D. Number:	574
Temperature	Readout I.D.:	-15HER/17-12
Diese Tibo	Took Check.	as V



Schematic of Traverse Point Layout

MAG	1 - CA 59	,	
Traverse		Stack	Cyclonic
Point	Head (Aps)	Temp.	Flow Check
Number	in. H <sub>2</sub> O	(T <sub>q</sub> ), °F	° from Null
REF.	POINT	L-3	
TEST	#1		
10:55	1.65	440	
11:05	1.65	441	
11:15	1.65	442	
11:25	1.65	441	
11:35	1.65	443	
TEST	*2		
12:19	1.65	444	
12:29	1.65	442	
12:39	1,65	444	
12:49	1.65	445	
12:59	1.65	444	
Average	1		

Traverse Point Number	Velocity Head ( $\Delta p_g$ ) in. H <sub>2</sub> O	Stack Temp. (T <sub>q</sub> ), °F	Cyclonic Flow Check ° from Null
TEST	<sup>#</sup> 3		
13:38	1.65	444	
13:48	1.65	445	
13:58	1.65	445	
14:08	1.65	444	
14:18	1.65	445	
	·		
			-
Average			

PACIFIC ENVIRONMENTAL SERVICES, INC. —



## HYDROCARBON SAMPLING FIELD DATA

Project No. F028

CLIENT: ONIZUKA	AFB		Date:	1-12-94
Sampling Location:		TURBINE :	#2 EXHAUST	
TANK # in :		108	117	10 G 6F99
FLOW METER; Time		Sample A "Ho"	Sample B	Sample C
•				
O	10:56	30.5 0.80	12:18 30.0 0.90	13:38 30,5 0.80
<u> </u>	11:06	<b>25.0</b> 0.90	12:28 25.2 0.98	13:48 24.8 0.90
	11:16	<u>18.6 o.88</u>	12:38 20.2 1.0	13:58 19.2 0.90
30	11:26	12.8 0.84	12:48 15.3 1.0	14:08 14.2 0.90
40	11:36	7.5 0.88	12:58 10.2 1.0	14:18 8.7 0.90
POST LEAK-CHECK				
0	12:22	7.6 0.00	13:40 10.4 0.00	14:38 8.5 0.00
	12:32	7.6 0.00	13:50 10.4 0.00	14:48 8,5 0,00
		OK	_OK_	OK
· · · · · · · · · · · · · · · · · · ·				
		TEST #1	TEST #2	TEST #3
				•
Pre Leak-checkOK			Post Leak-	check

PACIFIC ENVIRONMENTAL SERVICES, INC. —



Plant: <u>ONIZUKA AFB</u>

Date: <u>1-12-94</u>

Source/Sample Number: TURBINE #2 EXHAUST RUNS 1.2.3

1. 
$$Vm(std) = (17.64)(Vm)(Y) \left[ \frac{P_{bar} + (\triangle H/13.6)}{Tm} \right]$$

$$Vm(std) = (17.64)( )( )( )( )( ) + ( /13.6)$$

$$Vm(std) = NP \quad dscf.$$

2. Volume water vapor collected (standard conditions).

V(lo) = condensate from impingers and selica gel.

Vw(std) = (0.04707) V(10) = (0.04707)(

Vw(std) = NA scf.

3. Percent moisture, by volume.

Percent moisture, by volume.

$$BW_S = \frac{VW(std)}{VW(std) + VM(std)} = \frac{4.6\%}{() + ()} = \frac{4.6\%}{()} \times \frac{4.6\%}{()$$

4. Molecular weight, stack gas.

Dry molecular weight.

$$Md = 0.440(% CO_2) + 0.320(% O_2) + 0.280(% N_2 + % CO)$$

$$Md = 0.440 (2.5) + 0.320(17.5) + 0.280(80.0)$$

Md = 29./0 lb/lb-mole.

$$Ms = Md + Bw_s (18 - Md) = (29.10) + (0.046)(18 - 29.10)$$

Ms = 28.59 1b/1b-mole.



Plant: ONIZUKA AFB

Date: 1-12-94

Source/Sample Number: TURBINE #2 EXHAUST

5. Stack gas velocity average.

$$Vs(avg) = (85.49)(Cp)(V\Delta P) \left[ avg \sqrt{\frac{(Ts)}{(Ps)(Ms)}} \right]$$

$$Vs(avg) = (85.49)(1.00)(1.18) \sqrt{\frac{(460+445)}{(30.22)(28.59)}}$$

$$Vs(avg) = 103.2 \text{ ft/sec.}$$

6. Stack volumetric flow rate, actual conditions (stack temperature and pressure).

Qs = 
$$(60)(Vs)(A) = (60)(i03,2)(3./4)$$
  
Qs =  $19,450$  acfm.

7. Stack volumetric flow rate, standard conditions (68 degrees F, 29.92 Hg).

Q(std) = 
$$(17.64)(Qs)(1-Bw_s) \frac{Ps}{Ts}$$
  
Q(std) =  $(17.64)(19,450)(1-0.046) \frac{30.22}{905}$   
Q(std) =  $(17.64)(19,450)(1-0.046) \frac{905}{Ts}$ 

8. Isokinetic variation.

$$z_{I} = (K) \left[ \frac{(Ts)(Vm(std))}{(Ps)(Vs)(An)(\theta)(1 - Bw_{s})} \right]$$

$$z_{I} = (0.0945) \left[ \frac{()}{()} () () () (1 - )$$

FNY:P	E ME			•	OL A	\R								MOD	EL	T-1	021S-	21	
ENGINE MFR SOLAR GENERATOR MFR ELECTRIC MACHINERY									MODEL BEMAC II										
ENGINE SERIAL NUMBER  20646																			
	TURBINE /									AND GENERATOR								BOIL	ER
TIME	KW	RVAR	AMPS	٨	∙₽\$ 2	A	# 5 3	va	18	<b>T-7</b>		1-5	PCD	LOP	LOT	LO	TEMP	PSI	P
MDN																			
0100																			
0200																			:
0300						Ш					_								
0400						Ш					1								
0500											_		<b></b>						
0600											1								-
0700							Ш				$\perp$								ļ
0600							Ш										<u> </u>		
6900	<b>5</b> 50	-	15	7	4	1	5			GR.	_	οķ	73	57	1360	5	52	52	12
1000	550							2		B	) E	රිට	15	57	1360	<u>ড</u>	56	12.5	12
1100	కరు							ľ		(×8	2	K)	747	5+	136	^	40	12.5	15
1200	c					Ш	Ш				1		<u> </u>						
1300	550						Ш			3	2 (	ŚΜ	12	57	134	5	64	15.2	15
1400	22)					Ш				70;	1	10	72	57	136	5	66	13 ?	15
1500	540				Ш		Ш			214	2	105	12	58	1300	5	68	124	13
1000	540	<u> </u>				Ш				710	2	715	72	58	12/2	5	68	12.5	15
1700					<del>NVIII.</del>	Ļ	d To			_	_	-			1,			-	
1800		<u> </u>			Ш	Ш	Ш				_	20	4	re	1/1	<b>D</b> _	1/2:	15_	
1900						Ш	Ш			_	4		<u> </u>	_	<u> </u>	]	-		
2000					Ш	Щ	Щ				4		ļ		-	<del> </del>	<del>                                     </del>	<del> </del>	<del> </del>
2100		<u> </u>			Щ						4		<u> </u>	-	<del> </del>	<b>_</b>	-		<del> </del>
2200		<u> </u>				Ш	Щ			<b>.</b>	$\dashv$		<u> </u>	<del> </del>	<del> </del>	-	-	<del> </del>	-
2300	1	<u> </u>								Ļ			_	<u></u>		<u> </u>		1	<u>L</u>
<u>E</u>	NGINE	OPE	RATIN	G !	HÇK	JR.	5	+		LAD	JUE	U	+					-	-
PRE	PREVIOUS TOTAL 152497						7	MID	s			1							
TOTAL TODAY 10					- 1	DAY	l												
TOTAL TO DATE 152 50 7						- 8	W	GS											
REMARKS																			

PACKAGE NUMBER

1/12/94



# APPENDIX C LABORATORY REPORTS AND ANALYTICAL METHODS

- PACIFIC ENVIRONMENTAL SERVICES, INC. -

#### REPORT

## TRUESDAIL LABORATORIES, INC.

CHEMISTS - MICROBIOLOGISTS - ENGINEERS

DEVELOPMENT RESEARCH

PACIFIC ENVIRONMENTAL SERVICES, INC.

13100 Brooks Drive **CLIENT** 

Baldwin Park, CA 91706

Attn: S. Hugh Brown

SAMPLE 3 tanks from project PES/ONIZUKA AFB

DATE

Jan. 14, 1994 RECEIVED

14201 FRANKLIN AVENUE TUSTIN, CALIF. 92680

AREA CODE 714 • 730-6239

AREA CODE 213 • 225-1564

Jan. 18, 1994

FAX 714 • 730-6462

53691 LABORATORY NO.

INVESTIGATION

Total hydrocarbon analysis by SCAQMD method 25.2

#### **RESULTS**

The submitted samples were analyzed for  $CH_4$ , CO,  $CO_2$  and nonmethane hydrocarbons (as  $C_1$ ) by SCAQMD Method 25.2. Oxygen concentrations were determined by Orsat analysis.

The results obtained are as follows:

This report applies only to the sample or samples investigated and is not necessarily indicative of the quality or condition of apparently identical or similar products. As a mutual protection to clients the public and these Laboratories this report is submitted and accepted for the exclusive use of the client to whom it is addressed and upon the condition that it is not to be used, in whole or in part, in any advertising or publicity matter without references to the condition that it is not to be used, in whole or in part, in any advertising or publicity matter to those techniques to the condition of the condi without prior written authorization from these Laboratories.

#### PACIFIC ENVIRONMENTAL SERVICES, INC. LN 53691

SAMPLE	<u>ID</u>	NMVHC ppmvC <sub>1</sub>	CH <sub>4</sub>	CO ppmv	CO <sub>2</sub>	<b>0</b> <sub>2</sub> <u>%v</u>
Test-1	108	ND	7	60	18610	17.9
Test-2	117	11	6	58	18590	18.2
Test-3	106	ND	6	57	18700	18.1
Detection :	limit	4	2	2	2	0.2

ND = Not detect

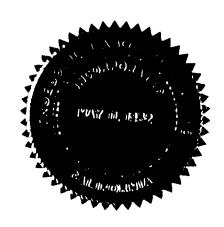
#### TRUESDAIL LABORATORIES, INC.

Prepared by:

Xuan Huong Dang Analytical Chemist

Air Pollution Testing

Charles M. Pigueroa Project Manager Air Pollution Testing



TCA-SOP Revision: Draft Date: 11/91

Prepared By: Page 1 of 10

#### DETERMINATION OF HYDROCARBON EMISSIONS BY TOTAL COMBUSTION ANALYSIS (TCA) METHOD

#### Table of Contents

- 1.0 Applicability
- 2.0 Equipment List
- 3.0 Preparation of sampling train
- 4.0 Sampling at test site 5.0 Analytical procedure
- 6.0 Gaseous sample analysis
- 7.0 Condensate trap recovery 8.0 Calculations
- 9.0 Diagrams

Hydrocarbon - SOP Revision: Draft Page 2 of 10

Method

This method is based on SCAQMD METHOD 25.1

#### 1.0 Summary of Method

This procedure uses a sampling train comprising a stainless steel probe and a freeze-out trap connected to an evacuated seven-liter tank via a Magnehelic pressure differential gauge. The trap is used for collecting the condensable organic matter while the non-condensable gases are being collected in the tank.

#### 2.0 Equipment list

- 2.1 7-liter tank with vacuum gauge
- 2.2 Stainless steel condensate trap and probe
- 2.3 Magnehelic gauge
- 2.4 Metal Dewar flask with dry ice
- 2.5 Tank holder

#### 3.0 Preparation of sampling train

- 3.1 Determine the number of tanks and traps required, and match only outlet tanks with outlet traps for sampling at an outlet location. Similarly, the principle applies to inlets.
- 3.2 Using a high-volume vacuum pump, evacuate the 7-liter tanks to a pressure of 1 Torr or less three times, filling the tanks to one atmosphere between evacuations.
- 3.3 After the third evacuation, turn off the valves and check for leaks by allowing the tanks to stand for at least 16 hours, after which period any leaks will become apparent by a change in the vacuum gauge readings.
- 3.4 For convenience in transporting as well as ease of handling at the test site, two tanks can be placed in a wooden holder.

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- 3.5 Set up each train in the following sequence: tank, Magnehelic gauge, trap, and probe. Be sure to determine the proper torque required to tighten these crucial connections, as too much torque will spoil the fittings for future conections while too little torque will result in leakage.
- 3.6 Perform a leak check on each train by introducing nitrogen into the connections at the probe tip. Magnehelic needle will deflect and should return to the same "zero" point after a short while. This signifies a good leak check.
- 3.7 The assembly is now ready for transport to the test site. It should be noted that triplicate samples are usually taken at the inlet locations and duplicates for outlets.

#### 4.0 Sampling at test site

- 4.1 Conduct pre-test flow and temperature measurements according to method described in Truesdail Laboratories, Inc. S.O.P. Before the stack traverse, make sure that both the air pollution control unit and the production line is running at the conditions in the test protocol or on the AQMD's "Permit to Construct."
- 4.2 Immerse the traps into the metal Dewar flask filled with crushed dry ice to a depth of four inches. Maintain the dry ice level for the whole duration of the test.
- 4.3 Uncap the probe tips and wrap the tips as close as possible to each other in order to satisfy duplicate or triplicate sampling. Avoid touching the tips to prevent contamination.
- 4.4 Insert the probes into the sampling port carefully so as not to scrape the port walls on the way in. Position the tips near the center of the duct. The port should then be sealed off with duct tape.
- 4.5 Coordinate all sampling stations, i.e., begin and end the test simultaneously. Integration times will vary from 30 to 120 minutes.
- 4.6 During the sampling period, record the vacuum gauge reading at 5-minute intervals and make field notes regarding any unusual events which may affect subsequent analytical results. Examples being: (1) the plugging of flow due to a frozen line, and (2) problems associated with the production line.

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- 4.7 Adjust the flow into the tanks such that between 5 to 10 inches of vacuum remain at the end of the sampling period. Close all valves at the end of the test.
- 4.8 Remove the probes from the duct carefully and cap off the probe tips.
- 4.9 Perform a post-test leak check by opening the valves. The Magnehelic needle will deflect and will settle down to its original position if there are no leaks. Make a note otherwise.
- 4.10 Label all tanks and traps accordingly, i.e., include the name of client, sampling date and location, and test and tank number.
- 4.11 For verification and quality control purposes, conduct a post-test flow and temperature measurement.
- 4.12 Transport the assemblies to the laboratory for analysis.
- 4.13 Disconnect the sampling trains and plug the open end of the traps. Store the traps in a dry ice or household freezer until they can be processed.
- 4.14 Measure the pressure in the tanks with a manometer and add pre-purified dry nitrogen to an absolute pressure of at least 860 Torr.
- 4.15 Record these pressures as they will be needed for further analytical computations. The corrected barometric pressure and temperature are to be recorded as well.

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#### 5.0 Analytical description

This involves the separate determinations of carbon monoxide, methane and carbon dioxide, and the combined determination of  $C_2$  and higher molecular weight hydrocarbons. Results are reported as parts per million (ppm) and pounds per hour (lb/hr) as carbon.

The gaseous portion and condensables portion are analyzed separately. The analysis of the gaseous portion requires a gas chromatographic column to separate and elute, in order, carbon monoxide, methane and carbon dioxide in the sample. The separated components are then methanized and detected by a flame ionization detector (FID). The amount of methane measured by the FID is recorded on a chromatogram strip chart.

The analysis of the volatile hydrocarbons in the gaseous portion requires a gas chromatographic column preceded by a 7-inch loop of Tenax material placed in an ice bath (0°C) which absorbs the C<sub>3</sub> and higher hydrocarbons. The C<sub>2</sub> hydrocarbons are separated from carbon monoxide, methane and carbon dioxide in the sample (which elute together) by the chromatographic column. After C<sub>2</sub> is eluted, the carrier gas flow direction is reversed and the Tenax loop is heated with boiling water (100°C) to desorb the remaining hydrocarbons. As each component is eluted, it passes through a catalytic oxidizer which converts it to carbon dioxide. Each carbon dioxide peak is measured by a non-dispersive infrared (NDIR) spectrophotometer, utilizing a carbon dioxide detector, and quantified by a computing integrator (GC/NDIR).

The condensable portion of the sample is analyzed for total hydrocarbons as carbon by volatilizing the trap contents and catalytically oxidizing everything to carbon dioxide which is then collected in an evacuated vessel and quantitatively determined by the FID as mentioned in 5.2.

In order to obtain meaningful analytical data, it is necessary to procure accurate reference standards and to calibrate the instruments with these standards at frequent intervals. Known concentrations of carbon monoxide, methane, carbon dioxide and propane in dry nitrogen are purchased from a vendor and are NBS traceable (+/-2%).

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#### 6.0 Gaseous sample analysis

- 6.1 Record the room temperature and barometric pressure.
- 6.2 After the sample tanks have settled down to room temperature, the absolute pressure is measured with a mercury manometer. The tanks are then pressurized with dry nitrogen to at least 32 inches of mercury (absolute), re-equilibrated, and measured again. The measurements are recorded and a dilution factor calculated.
- 6.3 The GC/FID instrument is calibrated with a standard gas sample.
- 6.4 The pressurized sample is used to flush the sample loop on the gas chromatograph with sample and the injection valve actuated to place the loop into the carrier circuit.
- 6.5 The sample fractions are eluted in order methane, carbon monoxide, and carbon dioxide and plotted on a strip chart chromatogram.
- 6.6 A computer instantaneously integrates and calculates each sample peak with the appropriate constants and correction factors, and reports these values at the end of the chromatogram. Replicate runs are made until ± 5% maximum deviation is obtained.
- 6.7 The GC/NDIR instrument is standardized with a standard gas sample.
- 6.8 The pressurized sample is used to flush the sample loop on the gas chromatograph with sample.
- 6.9 The injection valve is actuated, placing the sample loop into the carrier circuit with the Tenax loop immersed in an ice bath.
- 6.10 When the  $C_2$  hydrocarbons have been eluted or their retention time passed, the carrier gas flow through the Tenax and GC column is reversed and the Tenax loop immersed in boiling water. The  $C_3$  and higher molecular weight hydrocarbons are eluted together and plotted with the previous peaks on a strip chart chromatogram after detection by the NDIR analyzer.

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- 6.11 A computer instantaneously integrates and calculates each sample peak with the appropriate constants and correction factors and reports these values at the end of the chromatogram. Replicate runs are made until ± 5% maximum deviation is achieved.
- 6.12 In the event that any of the hydrocarbon peaks are high enough to make the analyzer off scale, the sample is reanalyzed using a smaller loop and an appropriate standard.

#### 7.0 Condensate trap recovery

- 7.1 Place the trap in a dry ice cooling bath and then heat the ends of the trap with a Bunsen burner to drive the hydrocarbons into the cold section of the trap away from the plugs.
- 7.2 After a minimum of five minutes in the cooling bath, the plugs are removed from the trap and the trap is then connected to the carrier gas on one end and a 1.8-liter evacuated vessel on the other end. The trap is purged at a rate similar to the sampling rate until at least 1 liter of purged gases are collected for at least six minutes.
- 7.3 Remove the purge gas vessel. Another evacuated collection vessel is the attached to the NDIR effluent and flow re-established in a push-pull fashion. The trap is connected to the oxidizer, followed by a sulfuric acid bubbler to remove moisture and a NDIR carbon dioxide analyzer to indicate how much hydrocarbon is left in the trap.
- 7.4 The cooling bath is removed and the trap is slowly heated with a Bunsen burner until the stainless steel trap reaches a dull red glow and the sample is eluted from the trap as indicated by the NDIR.
- 7.5 The collection vessel is analyzed for carbon dioxide by GC/FID as in 6.1 thru 6.6.
- 7.6 The purge gas vessel is analyzed for volatile hydrocarbons by GC/NDIR as in 6.7 thru 6.12.

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7.7 For traps where low condensables are expected, especially outlet ones, with the possibility of high moisture and carbon dioxide in the samples, the trap is placed in a water bath at room temperature and connected to a short loop of stainless steel (1/4-inch 0.D.) packed with quartz wool and placed in a dry ice bath. The purge step is then performed and the purge loop incorporated into the trap analysis as in 7.3. This alternative procedure will minimize the amount of unpurged stack carbon dioxide which may have dissolved in the trap condensate.

#### 8.0 Calculations

- 8.1 Condensable hydrocarbons:
  - (a) Integrate the area of the standard.
  - (b) Integrate the area of the sample.
  - (c) Calculate the concentration in ppm of carbon equivalent as follows:

$$C_{smp1} = \frac{C_{std} \times A_{smp1} \times V_{ves} \times \frac{P_{tank}}{29.9} \times \frac{520}{460 + T}}{A_{std} \times V_{tank} \times \frac{P_{tank}}{29.9} \times \frac{520}{460 + T}}$$

where  $C_{smpl}$  = concentration of the sample in ppm,  $C_{std}$  = concentration of the standard in ppm,  $A_{smpl}$  = area of the sample,  $A_{std}$  = area of the standard,  $V_{tank}$  = volume of the sample tank in liter,  $V_{ves}$  = volume of the collection vessel in liter,  $P_{tank}$  = pressure (absolute) of the tank in inches of mercury, and T = room temperature in  $^{\circ}F$ .

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- 8.2 Volatile hydrocarbons and gaseous components:
  - (a) Integrate the area of the standard.
  - (b) Integrate the area of the standard components.
  - (c) Calculate the concentration in ppm of carbon equivalent as follows:

$$C_{smpl} = \frac{C_{std} \times A_{smpl} \times \frac{P_B + P_f}{P_B + P_i}}{A_{std}}$$

where  $P_B$  = barometric pressure (net) in inches of mercury,

 $P_i$  = residual pressure of sample tank, and  $P_f$  = final pressure of sample tank after  $N_2$  addition.

8.3 The following formula is used for the computation of the emission rate in lb/hr carbon:

$$\frac{\textit{C x 12 lb/mole x } \textit{Q}_{sd}}{\textit{3.79 x 10}^{8} \textit{ ft}^{3}/\textit{lb.mole}}$$

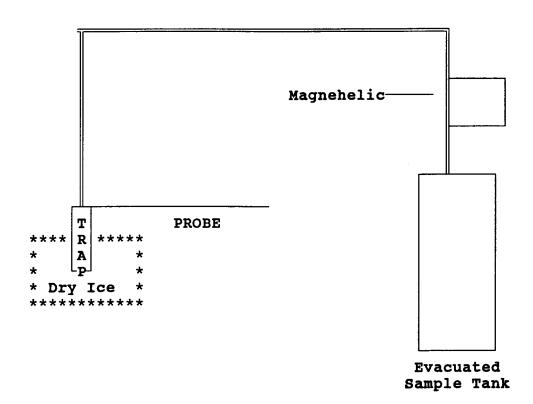
where C = total concentration of hydrocarbons present in ppm, and  $Q_{sd}$  = flow rate in standard ft<sup>3</sup>/hr (dry).

#### 9.0 Diagrams

Diagrams of the assembly is shown on the following page. Note that the distance between the dry ice top surface and the sampling probe of the trap should be at least one inch so that water will not freeze in the narrow sampling tube.

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#### TCA Sampling Train Setup



Revision: Draft

Date: 11/91
Prepared by: XHD
Approved by: PDM
QA Approval: TJP
Page: 1 of 8

#### STANDARD OPERATING PROCEDURE FOR: TOTAL HYDROCARBON ANALYSIS BY SCAQMD 25.1

#### Table of Contents

- 1.0 Method
- 2.0 Principle
  3.0 Applicability
  4.0 Apparatus
  5.0 Reagents
  6.0 Procedure
- 7.0 Calculations

25.1/SOP Revision: Draft Page: 2 of 8

#### STANDARD OPERATING PROCEDURE FOR: TOTAL HYDROCARBON ANALYSIS BY SCAQMD 25.1

#### Method

This method is based on SCAQMD METHOD 25.1. It is intended for the determination of total hydrocarbons in source emissions.

#### 1.0 Principle

Samples are collected in two fractions by using an evacuated tank to draw gases through a condensate trap chilled in dry ice. In this modification of the method, both fractions are separately analyzed for carbon monoxide, methane and carbon dioxide, in addition to a combined determination of C2 and higher molecular weight hydrocarbons. Results are reported as parts per million (PPM) in the sample, and as pounds of carbon per hour from the source.

#### Applicability

This method is applicable to determination of stationary source emmissions including incinerators, boilers and absorbers.

#### 2.0 Equipment Required

- 2.1 GC/FID
- 2.2 GC/NDIR
- 2.3 Ice water bath: 1000ml Beaker filled with ice and deionized water.
- 2.4 Boiling water bath: 1000 ml Beaker filled with deionized water, wire gauze, tripod and Bunsen burner.
- 2.5 Dry ice.
- 2.6 Evacuated 1.8 liter vessel.
- 2.7 Condensate trap

#### 3.0 Reagents Required

See 3.1 and 3.2 of SOP titled TCA 25.2/SOP

#### 4.0 Procedure

This method requires the use of two different instruments, GC/FID and GC/NDIR and Orsat equipment. The samples are collected in two fractions, a gaseous fraction and a condensible fraction. Each fraction must be analyzed separately using both instruments.

- 4.1 Preparation of Gaseous Fraction of Sample
  - 4.1.1 Record room temperature and barometric pressure.
  - 4.1.2 When the sample has equilibrated to room temperature, measure the absolute pressure in the sample cylinder with a mercury manometer, and record the result as residual vacuum (R.V.).
  - 4.1.3 Pressurize the sample cylinder with ultra-pure nitrogen to at least 6 inches of mercury (absolute), then allow to equilibrate to room temperature (about 5 minutes).
  - 4.1.4 Repeat the measurement of absolute pressure, and record the result as pressure of nitrogen (PN<sub>2</sub>).
  - 4.1.5 Calculation for dilution factor (XF) and sample amount (SA) (see 5.1).
- 4.2 Preparation of Condensable Fraction of Sample

Note: Inlet traps and outlet traps differ substantially in the level of hydrocarbons present, and so require slightly different handling. Several of the steps below will contain alternative instructions for onlet and outlet traps. Inlet traps and outlet traps are kept separate; inlet traps are not used to sample outlets, and vice versa in order to avoid sample carry-over problems.

- 4.2.1 Evacuate two 2 liter tanks to -30" Hg. Prepare a solution of 0.1N H<sub>2</sub>SO<sub>2</sub> for the knock-out which is connected between the catalyst and the NDIR. The water knock-out also requires an ice bath.
- 4.2.2 Determine the baseline for the NDIR. On range 3, the baseline will usually be between 10 and 15 at room temperature.

- 4.2.3 Remove the trap from the freezer. Inlet traps: immerse the trap in a dry ice bath, connect one side of the trap to the carrier gas, and the other to an evacuated tank. Outlet traps: connect one side of the trap to the carrier gas, and the other side to a cold finger which is immersed in a dry ice bath (the trap itself is allowed to warm to room temperature). The other side of the cold finger is connected to an evacuted tank.
- 4.2.4 Start the carrier gas flowing through the trap for 5 minutes, until the pressure the pressure in the tank is about -10" Hg. This tank is then analyzed and added to the sample collected in the trap.
- 4.2.5 Connect the trap to the catalyst column. Attach a second evacuated cylinder to the outlet of the NDIR. If a inlet is being burned, use an eight liter cylinder. If an inlet is being burned, use a two liter flask. Perform a leak check by turning on the carrier gas flow briefly. Watch the bubbles in the water knock-out. When the bubbles stop with the system pressurized and the carrier gas is turned off, there should be a complete equlibrium in the drop-out. The presence of water moving backwards in the drop-out signifies a leak in the trap or catylist. If bubbles continue to flow, there is a leak in the NDIR or tank connection.
- 4.2.6 Start the carrier gas flow, remove the dry ice bath and begin heating the trap with the burners. For inlet traps, set the supplemental oxygen flow to match the carrier gas flow. For outlet traps, burn the cold finger along with the trap: supplemental oxygen is not needed in most cases. Observe the reading on the NDIR. It should quickly begin to rise.
- 4.2.7 Continue burning the trap until the NDIR reading falls to near the blank value. The trap is allowed to cool to room temperature, and then is briefly heated again to determine whether all the condensable hydrocarbons have been driven off. If the NDIR reading does not increase during this final check the trap is clean. It is important that hydrocarbons will condense in the tubing and

that it must be burned out with the trap. Also any place there is a connector, it must be heated to make sure that the inside is cleaned.

- 4.2.8 The tank from the burn and the purge are pressurized and analyzed (see above for gas sample portion).
- 4.3 GC/FID Analysis of Samples
  - 4.3.1 Check carrier gas (UP He), if low, change the gas tank.
  - 4.3.2 Change the output from "test" to "1", change the range from "Bal" to "10".
  - 4.3.3 Increase He to 50 psi.
  - 4.3.4 Turn on H2 to 30 psi.
  - 4.3.5 Ignite the FID.
  - 4.3.6 Turn on Air to 20 psi.
  - 4.3.7 Increase He to 80 psi.
  - 4.3.8 Turn FID Zero Suppression on, and switch detector Output from "1" to "2".
  - 4.3.9 Let the instrument warm up for 20 minutes.
  - 4.3.10 Start the Maxima program of the Dynamic Solutions system, and set up the automatic data acquisition according to the manual Maxima 820.
  - 4.3.11 Standard Calibration Calibration standards are commercial stock standard mixtures made from certified gas company.
    - (a) Inject a standard sample through a sample loop on the GC.
    - (b) Press "Run" on the GC to place the loop into the carrier gas circuit.
    - (c) The calibration standards are run in replicate until +/- 2% maximum deviation is obtained.
  - 4.3.12 The sample is injected in the same manner as the standard, steps 4.3.11: a, b, and c.

- 4.4 GC/NDIR Analysis of Samples
  - 4.4.1 Turn on the instrument and allow the GC oven to reach analysis temperature (approximately about 800 F).
  - 4.4.2 Start Nitrogen flow.
  - 4.4.3 Turn on the integrator: change the RANGE from "Tune" to "1", set GAIN from "0" to "5".
  - 4.4.4 When the GC oven reaches 800 F, turn the ZERO tune all the way to the right, and then adjust the SOURCE BALABCE until it reads between 35-40.
  - 4.4.5 Adjust the ZERO tune until the display reads "2" by turning it to the left.
  - 4.4.6 Equilibrate the Tenax loop in the ice water bath.
  - 4.4.7 Run a blank sample to check for carry-over prior to running standard.
  - 4.4.8 Standard Calibration Calibration standards are commercial stock standard from certified gas company.
    - (a) Flush the sample loop with the pressurized standard, then switch the injection valve to inject the sample.
    - (b) Press "Start" on the integrator.
    - (c) When the  $\mathrm{C}_2$  has been eluted or their expected retention time has passed, the carrier gas flow direction is reversed, and the Tenax loop is immersed in boiling water to desorb the remaining hydrocarbons.
    - (d) The calibration standards are repeated until +/- 5% maximum deviation is obtained.
  - 4.4.9 Inject the sample in the same manner as the standard, steps 4.4.8: a, b, and c
  - 4.4.10 Repeat the analysis of samples until +/-5% maximum deviation is obtained. If any of the hydrocarbons peaks are off scale, recalibrate the GC using a new standard that is closer to the concentration of the sample and then reanalyze the sample.

4.5 Dilution

See 4.5 of the SOP titled TCA 25.2

4.6 Orsat analysis of samples

See SOP titled 7.1

#### 5.0 Calculations

- Condensable Hydrocarbons
  - The intergrated area for the standard and the sample are available directly from the chart recorder.
  - Calculate the concentration in ppm of carbon 5.1.2 equivalent as follows:

$$C_{smp1} = \frac{C_{std} \times A_{smp1} \times V_{ves} \times \frac{P_{tank}}{29.9} \times \frac{520}{460 + T}}{A_{std} \times V_{tank} \times \frac{P_{tank}}{29.9} \times \frac{520}{460 + T}}$$

where

 $C_{smpl} = concentration of the sample in ppm,$ 

 $C_{\text{std}}^{\text{smpl}} = \text{concentration of the standard in ppm,}$   $A_{\text{smpl}}^{\text{smpl}} = \text{area of the sample,}$ 

 $A_{\text{std}} = \text{area of the standard,}$   $V_{\text{tank}} = \text{volume of the sample tank in liter,}$   $V_{\text{ves}} = \text{volume of the collection vessel in liter,}$   $P_{\text{tank}} = \text{pressure (absolute) of the tank in inches of}$ mercury,

= room temperature in F.

- Volatile hydrocarbons and gaseous components 5.2
  - The integrated area for the standard and the sample are available directly from the chart recorder.
  - 5.2.2 Calculate the concentration in ppm of carbon equivalent as follows:

$$C_{smpl} = \frac{C_{std} \times A_{smpl} \times \frac{(P_B + P_f)}{(P_B + P_i)}}{A_{std}}$$

where:

 $P_B$  = barometric pressure (net) in inches of mercury,

P<sub>i</sub> = residual pressure of sample tank,

P<sub>f</sub> = final pressure of sample tank after N2 addition.

The following formula is used for the computation of the emission rate in lb/hr carbon:

$$\frac{\textit{C} \times 12 \textit{lb/mole} \times \textit{Q}_{\textit{sd}}}{3.79 \times 10^{8} \textit{ft}^{3} / \textit{lb.mole}}$$

#### where:

C = total concentration of hydrocarbons
 present in ppm,

 $Q_{sd}$  = flow rate in standard cubic feet per hour (dry).



APPENDIX D
CALIBRATION DATA

----- PACIFIC ENVIRONMENTAL SERVICES, INC. --



2600 CAJON BOULEVARD, SAN BERNARDINO, CA 92411

(909) 887-2571 FAX. (909) 887-0549

#### CERTIFICATE OF ANALYSIS: EPA PROTOCOL GAS RECERTIFICATION

Customer

PACIFIC ENVIRONMENT SER ATTN: STEVE HERNANDEZ 13100 BROOKS DRIVE BALDWIN PARK CA91706

Assay Laboratory Scott Specialty Gases 2600 Cajon Boulevard San Bernardino, CA 92411

Purchase Order 0640-102 **Scott Project #** 25837.001

ANALYTICAL INFORMATION

Certified to exceed the minimum specifications of EPA Protocol 1 Procedure #G1, Section Number 3.0.4

Cylinder Number

ALM027046

Certification Date

12-23-93

Acid Rain Exp.

Cylinder Pressure

2000PSIG

**Previous Certification Dates** 

05-20-93 General Exp. 12-23-95

ANALYZED CYLINDER

Components NITRIC OXIDE Certified Concentration

43.83PPM

Analytical Uncertainty\* ±1% NIST Traceable

Balance Gas: Nitrogen

NOX

44.17PPM

\*Analytical uncertainty is inclusive of usual known error sources which at least includes reference standard error & precision of the measurement processes

REFERENCE STANDARD

GMIS =

**Expiration Date** 

09-94

Cylinder Number

ALM033883

Concentration

100.4PPM

INSTRUMENTATION

Instrument/Model/Serial # TECO / 10AR-38644-258

Last Date Calibrated

10-29-93

**Analytical Principle** 

Chemi-Luminescent

#### ANALYZER READINGS (Z=Zero Gas R=Reference Gas T=Test Gas r=Correlation Coefficient)

#### Components NITRIĆ OXIDE

#### **Previous Certification**

Date: 05-20-93	Response Units: mv		
Z1=	R1=	T1=	
R2=	<b>72</b> =	T2=	
<b>Z3</b> =	T3=	R3=	
Avg. Conc. of C	ust Cyl.	43.83PPM	

#### Third Triad Analysis

Date: 12-23-93		Response Units: mv		
Z1=	0.00	R1=	97.0	T1= 41.8
R2=		<b>Z2=</b>	0.00	T2= 41.8
<b>Z3</b> =		T3=		<b>R3=</b> 97.0
Avg.	Conc. of C	ust Cy	i.	43.83PPM

#### Calibration Curve

Concentration=	Ax + B
A =1.0083945	
B =-0.139473	

Date:	Respo	mse Units:
Z1=	R1=	T1=
R2= Z3=	<b>72=</b>	T2=
<b>Z3</b> =	T3=	R3=
Avg. Conc. o	f Cust Cyl.	

Dute:	Response Units:			
Z1=	R1=	T1=		
R2=	<b>72</b> =	T2=		
<b>Z3</b> =	T3=	R3==		
Avg. Conc.	of Cust Cyl.			

Concentration=				
•				

Date:	Respo	ense Units:
Date: Z1= R2= Z3=	R1=	T1=
R2=	72=	T2=
Z3=	T3=	R3=
	of Coast Carl	

Date:	Respon	nse Units:
Z1=	R1=	Ti=
R2=	7.2=	T2=
<b>73</b> =	T3=	R3=

Concentration=					

Avg. Conc. of Cust Cyi.

Avg. Conc. of Cust Cyi.

SPECIAL NOTES: IF THIS PRODUCT IS USED FOR ACID RAIN COMPLIANCE, THE ACID RAIN DATE NOTED ABOVE APPLIES PER 40 CFT PARTYS, APPENDIX H. OTHERWISE THE GENERAL EXPIRATION DATE APLLIES.

Analyst



2600 CAJON BOULEVARD, SAN BERNARDINO, CA 92411

(909) 887-2571 FAX: (909) 887-0549

#### CERTIFICATE OF ANALYSIS: EPA PROTOCOL GAS

Customer:

Pacific Environmental Services 13100 Brooks Drive Baldwin Park, CA 91706-0740

Assay Laboratory

Scott Specialty Gases 2600 Cajon Boulevard San Bernardino, CA 92411 Purchase Order 0640-102 Scott Project # 24700

#### **ANALYTICAL INFORMATION**

Certified to exceed the minimum specifications of EPA Protocol 1 Procedure #G1, Section Number 3.0.4

Cylinder Number

ALM027851

**Certification Date** 

03-17-93

**Expiration Date** 

Cylinder Pressure

1950 psig

**Previous Certification Dates** 

NONE

09-13-94

ANALYZED CYLINDER

Components NItric Oxide **Certified Concentration** 

Analytical Uncertainty\* ± 1 % NIST Traceable

22.51 ppm

NOX

Balance Gas: Nitrogen

22.82 ppm

\*Analytical uncertainty is inclusive of usual known error sources which at least includes reference standard error & precision of the measurement processes

COLUMN S REFERENCE STANDARD

Type \* **SRM 2629A**  **Expiration Date** 

09-93

Cylinder Number FF28519

Concentration

19.3 ppm

77.12

0.000

90.50

T3=

T1 = 90.20

T2= 90.54

R3= 77.21

22.60 ppm

INSTRUMENTATION Instrument/Model/Serial # TECO / 10AR / 14853-150

Last Date Calibrated

Z1=

R2=

73-

1-20-93

**Analytical Principle** Chemi-Luminescent

#### ANALYZER READINGS (Z=Zero Gas R=Reference Gas T=Test Gas r=Correlation Coefficient)

#### Components

Nitric Oxide

#### First Triad Analysis

Date: 03-01-93

Second Triad Analysis Date: 03-17-93 Response Units: mv

0.000

77.24

0.000

Avg. Conc. of Cust Cyl.

Calibration Curve

 $Ax^2+Bx+C$ 

Concentration= A =-0 000504296 B=1.0489880 C =-0.292506

Dota		R	CADORAC	Units: 1	nv
Avg. Conc. of Cust Cyl. 22.41 ppm					
Z3=	0.000	T3=	90.94	R3=	78.28
R2=	0.000 78.30 0.000	<b>Z2=</b>	0.000	T2=	90.92
Z1=	0.000	R1=	78.33	T1=	90.95

Response Units: mv

R1= 78.33 T1= 90.95

Date:	Respo	Response Units: mv		Respo	nse Units: mv
Z1=	R1=	T1=	Z1=	R1=	T1=
R2=	<b>Z2</b> =	T2=	R2=	<b>72</b> =	T2=
<b>Z3</b> =	T3=	R3=	Z3=	T3=	R3=
Avg. Conc. of Cust Cyl.		Avg. Conc.	of Cust Cyl.		

ı	Concentration=
l	, ,
- 1	

Date:	Respo	nse Units:
Date: Z1= R2= Z3=	R1=	T1=
R2=	<b>72=</b>	T2=
Z3=	T3=	R3=
	of Cust Cyl.	

Date:	Response Units:	
Z1=	R1=	T1=
Z1= R2= Z3=	<b>72</b> =	T2=
Z3=	T3=	R3=
	c. of Cust Cyl.	

Concentration=

Special Notes



2600 CAJON BOULEVARD, SAN BERNARDINO, CA 92411

(909) 887-2571 FAX: (909) 887-0549

#### **CERTIFICATE OF ANALYSIS: EPA PROTOCOL GAS**

PACIFIC ENVIRONMENTAL SER. PO# 0640-102 13100 BROOKS AVE BALDWIN PARK, CA 91706

Assay Laboratory Scott Specialty Gases 2600 Cajon Boulevard San Bernardino, CA 92411

Purchase Order 0640-102 Project # 27964.001

**ANALYTICAL INFORMATION** 

Certified to exceed the minimum specifications of EPA Protocol 1 Procedure #G1, Section Number 3.0.4

Cylinder Number Cylinder Pressure ALM034155

**Certification Date** 

10-06-93

GENERAL Date 10-06-95

ACID RAIN DATE

ANALYZED CYLINDER

Components NITRIC OXIDE Certified Concentration

22.90 PPM

Analytical Uncertainty\* ±1 % NIST Traceable

Balance Gas: Nitrogen

NOX

.....

23.46 PPM

\*Analytical uncertainty is inclusive of usual known error sources which at least includes reference standard error & precision of the measurement processes

REFERENCE STANDARD

GMIS

**Expiration Date** 

Cylinder Number

ALM033911

Concentration

· 24.97 ppm

**INSTRUMENTATION** 

Instrument/Model/Serial#

TECO / 10AR / 38644-258

**Last Date Calibrated** 

07-26-93

**Analytical Principle** 

Chemi-Luminescent

ANALYZER READINGS (Z=Zero Gas R=Reference Gas T=Test Gas r=Correlation Coefficient)

Components

First Triad Analysis

Second Triad Analysis

**Calibration Curve** 

Nitric Oxide

Date	09-29-93	Resp	onse Units: mv
Z1=	0.00	R1= 96.4	4 T1= 88.5
R2= Z3=	96.4	<b>22=</b> 0.00	0 12= 88.5
<b>Z3</b> =	0.00	T3= 88.5	s R3= 96.4
Avo.	Conc. of C	ust CvL	22.91 poza

Date:	10-06-93	R	Response Units: mv	
Z1=	0.00	R1=	96.4	T1= 88.4
R2==	96.4	<b>Z2=</b>	0.00	T2= 88.3
<b>Z3</b> -	0.00	T3=	88.3	R3= 96.4
Avg. (	Conc. of C	ust Cy	L.	22.88 ppm

Concentration=	Ax + B
A=1.001	
B =0.05283	
ŀ	

Date:	Respo	onse Units: mv
Z1=	R1=	T1=
R2=	<b>Z2=</b>	T2=
<b>Z3=</b>	T3=	R3==
Avg. Conc	of Cust Cyl.	

Date:	Response Units: mv	
Z1=	R1=	T1=
R2=	<b>Z2=</b>	T2=
Z3=	T3=	R3=
Avg. Conc.	of Cust Cyl.	

Concentration=	-	

Date:	Response Units: mv	
Date: Z1= R2= Z3=	R1=	Ti=
R2=	<b>Z2</b> =	T2=
Z3=	T3=	R3=
Avg. Conc.	. of Cust Cyl.	

Date:	Response Units: mv	
Z1=	R1=	T1=
R2=	Z2=	T2=
<b>Z3=</b>	T3=	R3=

Concentration=	

SPECIAL NOTES: IF THIS PRODUCT IS USED FOR ACID RAIN COMPLIANCE, THE ACID RAIN EXPIRATION DATE NOTED ABOVE APPLIES PER 40 CFT PART 75, APPENDIX H. OTHERWISE THE GENERAL EXPIRATION DATE APPLIES. ANALYST Thomas & Wilson



Shipped From: 2600 CAJON BLVD.

SAN BERNARDINO

CA 92411

Phone: 909-887-2571

Fax: 909-887-0549

#### CERTIFICATE OF ANALYSIS

PACIFIC ENVIRONMENTAL SER

F'0# 0640-102

13100 BROOKS DRIVE

PROJECT #: 02-28622-001

PO#: 0640-102

ITEM #: 02024520 4AL

DATE: 11/11/93

BALDWIN PARK

CA 91706

CYLINDER #: ALMO35757

ANALYTICAL ACCURACY: +/-1%NIST

FILL PRESSURE: 2000PSIG

BLEND TYPE : CERTIFIED MASTER GAS

ANALYSIS REQUESTED GAS \_(MOLES) \_\_CONC\_MOLES\_ COMPONENT 11.09 F'CT CARBON DIOXIDE 11. F'CT 50. F'F'M F'F'M 50.11 CARBON MONOXIDE 11. FCT OXYGEN F'CT 11.00 **NITROGEN** BAL BAL

2000PSIG BIN#2 11-12-93 CRM1675 14.08%CO2 ALMO01136 CRM1678 47.2PPM AAL5970 CO CRM2659 20.63% ALMO17555 D2

ANALYST :/

PLUMSTEADVILLE, PENNŚYLVANIA / TROY. MICHIGAN / HOUSTON, TEXAS / DURHAM, NORTH CAROLINA SOUTH PLAINFIELD, NEW JERSEY / FREMONT. CALIFORNIA / WAKEFIELD. MASSACHUSETTS / LONGMONT. COLORADO BATON ROUGE. LOUISIANA



Shipped

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SAN BERNARDINO

CA 92411

Phone: 909-887-2571

Fax: 909-887-0549

#### CERTIFICATE OF ANALYSIS

PACIFIC ENVIRONMENTAL SER

STEVE HERNANDEZ

13100 BROOKS DRIVE

PROJECT #: 02-27898-001

PO#: 0640-102

ITEM #: 02024520

DATE:10/01/93

BALDWIN PARK

CA 91706

CYLINDER #: ALMO36879

ANALYTICAL ACCURACY: +-1%

FILL PRESSURE: 2000 PSIG

BLEND TYPE : ACUBLEND MASTER GAS

COMPONENT
CARBON DIOXIDE
CARBON MONOXIDE
OXYGEN
NITROGEN

TED GAS	ANALTS	212
MOLES_	_(MOL	ES)_
PCT	18.00	PCT
F'F'M	75.00	FFM
PCT	19.00	FCT
BAL.		BAL
	MOLES PCT PPM PCT	MOLES         (MOLES           PCT         18.00           PFM         75.00           PCT         19.00

2000 PSIG BIN#2 10-08-93 CRM1679 ALM10524 CD CRM1675 ALMO01136 CO2, CRM2659 ALMO17555 O2

ANALYST:

PLUMSTEADVILLE, PENNSYLVANIA / TROY, MICHIGAN / HOUSTON, TEXAS / DURHAM, NORTH CAROLINA SOUTH PLAINFIELD, NEW JERSEY / FREMONT, CALIFORNIA / WAKEFIELD, MASSACHUSETTS / LONGMONT, COLORADO BATON ROUGE, LOUISIANA



Shipped From:

2600 CAJON BLVD.

SAN BERNARDINO

CA 92411

Phone: 909-887-2571

Fax: 909-887-0549

CERTIFICATE OF ANALYSIS

PACIFIC ENVIRONMENTAL

STEVE HERNANDEZ

C/O FEDERAL EXPRESS

520 LAWERNCE EXPRESS WAY

SUNNYVALE

NOX

CA 94086

PROJECT #: 02-28936-001

PO#: F028-000

ITEM #: 02022913 4AL

DATE: 12/02/93

CYLINDER #: ALMO10841

ANALYTICAL ACCURACY: +/-1%NIST

FILL PRESSURE: 2000 PSIG

BLEND TYPE : CERTIFIED MASTER GAS

<u>COMPONENT</u> NITRIC OXIDE NITROGEN - OXYGEN FREE CONC MOLES

10. PPM
BAL

ANALYSIS
(MOLES)
10.36 PP

PPM BAL

MIROGEN - OXIGEN FREE

10.39 PPM

2000 PSIG BIN #1 12-01 GRAVIMETRIC MASTER GAS AGAINST NIST CERTIFIED WEIGHT INDEPENDENT LABORATORY

CERTIFIED TO HAVE BEEN BLENDED AND VERIFIED TO BE CORRECT BY ANALYSIS.

ANALYST:

PLUMSTEADVILLE, PENNSYLVANIA / TROY, MICHIGAN / HOUSTON, TEXAS / DURHAM, NORTH CAROLINA SOUTH PLAINFIELD, NEW JERSEY / FREMONT, CALIFORNIA / WAKEFIELD, MASSACHUSETTS / LONGMONT, COLORADO BATON ROUGE, LOUISIANA EVER READY THERMOMETER CO., INC. 401 PARK AVENUE SOUTH NEW YORK, NY 10016

#### REPORT OF CALIBRATION

LIQUID-IN-GLASS-THERMOMETER

CALIBRATED BY EVER READY THERMOMETER CO.

MARKED: ERTCO 611-3FC S/N-2269

RANGE: -20 TO +110 DEGREES C IN 1 DEGREE GRADUATIONS.

THERMOMETER CORRECTION READING (ITS-90)\*\*

0.0 C 0.0 C 37.0 -0.1 56.0 0.0

\*\* ALL TEMPERATURES IN THIS REPORT ARE BASED ON THE INTERNATIONAL TEMPERATURE SCALE OF 1990 (ITS-90) PUBLISHED IN THE METROLGIA 27, NO. 1, 3/10/90.

THIS THERMOMETER WAS CALIBRATED AGAINST A STANDARD CALIBRATED AT THE NATIONAL INSTITUTE OF STANDARDS AND TECHNOLOGY (NIST) FORMERLY THE NATIONAL BUREAU OF STANDARDS (NBS).

FOR A DISCUSION OF ACCURACIES ATTAINABLE WITH SUCH THERMOMETERS SEE NBS MONOGRAPH 150.

IF NO SIGN IS GIVEN ON THE CORRECTION, THE TRUE TEMPERATURE IS HIGHER THAN THE INDICATED TEMPERATURE; IF THE SIGN GIVEN IS NEGATIVE, THE TRUE TEMPERATURE IS LOWER THAN THE INDICATED TEMPERATURE. TO USE THE CORRECTIONS PROPERLY, REFERENCE SHOULD BE MADE TO THE NOTES GIVEN BELOW.

THE THERMOMETER WAS TESTED IN A LARGE, CLOSED-TOP, ELECTRICALLY HEATED, LIQUID BATH, BEING "IMMERSED" 76MM. THE TEMPERATURE OF THE ROOM WAS ABOUT 25 DEGREES C (77 DEGREES F). IF THE THERMOMETER IS USED UNDER CONDITIONS WHICH WOULD CAUSE THE AVERAGE TEMPERATURE OF THE EMERGENT LIQUID COLUMN TO DIFFER MARKEDLY FROM THAT PREVAILING IN THE TEST, APPRECIABLE DIFFERENCES IN THE INDICATIONS OF THE THERMOMETER WOULD RESULT.

THE TABULATED CORRECTIONS APPLY PROVIDED THE ICE POINT READING IS 0.0 DEGREES C. IF THE ICE-POINT READING IS FOUND TO BE HIGHER (OR LOWER) THAN STATED, ALL OTHER READINGS WILL BE HIGHER (OR LOWER) TO THE SAME EXTENT.

TEST NUMBER: 140381
DATE: 06/21/90
STANDARD SERIAL NO. 128239
NIST IDENTIFICATION NO. 88024

EVER READY THERMOMETER CO.



#### METER BOX TEMPERATURE READOUT CALIBRATION

CALIBRATED BY: Falset Algunger

DATE: 06-16-93

DATE: 06-16-93

	Inlet (°F)	Outlet (°F)	Thermometer (°F)
BOX 1A	32	32	32 ( 0°C)
	77	76	77 ( 25°C)
	212	211	212 (100°C)
BOX 2A	32	32	32 ( 0°C)
	76	76	77 ( 25°C)
	213	213	212 (100°C)
BOX 3A	28	27	32 ( 0°C)
	75	75	77 ( 25°C)
	213	213	212 (100°C)

NOTE:

Thermometer Standard Serial Number: 128239

Thermometer NIST I.D. Number: 88024

PACIFIC ENVIRONMENTAL SERVICES, INC. —



#### THERMOCOUPLE CALIBRATION

CALIBRATED BY: Robint Aiguren

DATE: 06-16-93

Thermocouple number	Thermocouple reading (°C)	Thermometer reading (°C)
TC-1	0.0 26.5 100.0	0.0 26.5 100.0
TC-2	OUT OF SERV	/ICE
TC-3	0.0 27.5 99.0	0.0 26.5 100.0
TC-4	OUT OF SERV	JICE
TC-5	0.0 26.6 99.0	0.0 26.5 100.0
TC-6	0.0 26.5 100.0	0.0 26.0 100.0
TC-7	0.0 26.2 100.0	0.0 26.0 100.0
TC-8	0.0 26.3 100.0	0.0 26.0 100.0
TC-9	0.0 26.6 100.0	0.0 26.0 100.0

PACIFIC ENVIRONMENTAL SERVICES, INC. —

5.2	J 260	· Alta
		$\Box$
<u> </u>	<i>)</i>	ناك

TC-10	0.0 26.3 100.0	0.0 26.0 100.0
TC-11	0.0 26.0 100.0	0.0 26.0 100.0
TC-12	0.0 26.0 100.5	0.0 25.0 100.0
C-1	NOT AVAILA	ABLE
C-2	NOT AVAILA	ABLE
S-1A	0.0 25.8 99.0	0.0 26.0 100.0
S-2A	0.0 25.5 99.0	0.0 26.0 100.0
S-16A	0.0 25.7 99.0	0.0 26.0 100.0
S-17A	0.0 25.9 99.0	0.0 26.0 100.0

Thermometer Standard Serial Number: 128239

Thermometer NIST I.D. Number: 88024

PACIFIC ENVIRONMENTAL SERVICES, INC.



#### DIGITAL THERMOMETER CALIBRATION

CALIBRATTED BY: Labout Nigurer

DATE: 06-16-93

Digital Thermometer	(°F)	Thermometer (°C)
Fisher	32.0 80.0 210.0	0.0 (32 °F) 26.5 (80 °F) 99.0 (210 °F)
Digital Thermometer	(°C)	Thermometer (°C)
Omega	1.0 26.0 98.0	0.0 26.5 98.0
Digital Thermometer	(°C)	Thermometer (°C)
Fluke (T1)	1.0 27.5 99.0	0.0 26.5 98.0
Digital Thermometer	(°C)	Thermometer (°C)
Fluke (T2)	1.1 27.4 99.0	0.0 28.0 98.0

NOTE:

Thermometer Standard Serial Number: 128239

Thermometer NIST I.D. 88024

PACIFIC ENVIRONMENTAL SERVICES, INC. —

DATE: DRY GAS METER IDENTIFICATION: 25507 Model 5-190 OPERATOR

WIN IDENTIFICATION: AL-20

 $\int$  PACIFIC ENVIRONMENTAL SERVICES, INC.

BAROMETRIC PRESSURE (Pb):\_\_\_ ( Note: Pb was selling during calibration) 29,55 in. Hg

APPROXIMATE	WET TEST	DRY GAS		TEMPER	TEMPERATURES		DRY GAS	TIME	FLOW RATE	METER	AVERAGE METER
CON ANIE	GAS VOLUME	METER VOLUME	MET TEST	_	DRY GAS METER	ER	PRESSURE (AP)	min.	cfm	COEFFICIENT (Yd.)	(Y <sub>d</sub> )
	Ť.,	**	3	INLET	OUTLET	AVERAGE	in H <sub>2</sub> O			5	
	······································		•	<b>:</b> :3	ژ. ژ.	(Ī <sub>d</sub> )					
0 40 A	0.000	384.603 0003	74 76	1	77.1	77	-1,35	13.0	0.407	1.008	
	T	290.028	777		7.7	١	-1.35			,	
<i>E</i> 3		295,427	74	1	77.4	77	-1.35	13.0	0,405	800%	
·		295.575	7.6		77.4		-1,35	75/	7 1100	, 020	1.008
		300,968	74	1	77,4	\ \ \	-1.35	10.0	0,405	1,007	
0.60	2,000	301.605	74 74	1	77.4	77	-2.35	12.0	0.606	1.010	
P1 = 29.48 -> B		309, 187	42	,	77.4	77	-2.35	12.0	0.603	1 1009	
	Т	3/6,689	74		77.4		~ 2,35	,	2 221		•
6		324.140	74	1	78.2	18	-2.35	14.0	0.604	1.012	
0 80 A	0.000	325.721	75	1	78,2	78	-3.65	/&.O	0,822	1.010	
	Т	335,979	74	,	78,2		-3.65	,	2 62 1		
Pr: 29.46 + B	Γ	346.174	74		78,2	78	-3,65	12.0	0.822	1.010	
	0,000	346,279	74 77	)	78, Z 78, Z	78	-3.65	/ね. 0	0.825	1.009	1.010
1 00	A 0,000	357.93/	7,4	. 1	78, 2	75	515	10.0	1.008	1.009	
	Т	369, 142	75		78,2		-5.15	,	, , , ,	. 230	
-	10.358	379,598	74		78.2	9.7	-5,15	10.0	1.00	,	
	C 10.325	390,237	77 74	1	78,2	78	-5.15	10.0	1.005	1.009	1.009
1.20	0,000	391.437	75	′	2,35	32	-7,10	9,0	1,203	1,008	
	0.000	402.922	42		28,2	70	-7.10	9	12011	/ // /	
	11,102	717.400	7,7		7.8/	1	2/10		1	00	/ ///
	C 11.100	425.608	74	J	18. Ý	78	-7.10	9.0	1,201	1.001	1,001

 $Q = 17.65 \frac{V_s}{\Theta} \frac{P_b}{(T_s + 460)}$  $Y_{ds} = \frac{V_{\star}}{V_{dg}} \times \frac{(\bar{I}_d + 460)}{(I_s + 460)} \times \frac{P_b}{(P_b + \frac{\Delta P}{4})}$ 

# DRY GAS METER AND ORTFICE CALIBRATION

1	Barometric Pressure		Dry Gas Meter No.
1	29.8		1557640
	Calibration Date 12-28-93 Calibrated by:		Meter Box No.
	12-28-93		Ħ
	Calibrated by:		Reference Dry Gas Meter No. 25507 Rockw
	1	ı	s Meter No. 2
18.11.12.	M		7507 Rockwell

ар роска рос 	.,			He I Gri
0	2.0	1.0	0.5	Orifice I Manometer I Setting I H="H2O I
792.980	782.267	771.657	761.348	Initial Reference DOM Reading V=ft3
1 086	267 I	557 [	48	and haved breed breed breed breed
802.990	792.290	781,789 1	771.359 1	Final I Reference I DGM I Reading I V=ft3 I
10.01	10.023	10.132	10.011	Reference Gas Volume Vr=ft3
. = .	23 1	23 -	-	ng pang pang pang pang
730.982 [	719.945	709.045 I	698.576 [	Initial I DGM I Reading I V=ft3 I
741.358	730.205	719,423	708.743	Final DGM Reading V=ft3
- <del></del> -		¤	_ =	and private private private private
10.376	10.260	10.378	10.167	Test DGM Volume Vd=ft3
74	I 74	76	I 76	Temperal
4 Port F				Temper -[
101	97 I	123	8   1	Temperature  ox DGM I Bo Inlet I Ou t=F I
87	84	87	83	THE WAR
74	91	87	84	I td I td
-O	13.4	18.7	25.6 I	3 7 7 W
	!	i i	1	Flow Rate O=cfm
0,991 1 1.931	0.74   1.003   1.968	0.53 1 0.994 1 1.903	0.38   0.998   1.837	50 AB
-0	1 1.96	1 1,90	1 1.83	He ta

	AVERAGE	
~		 
	0.997	 
		 ,
-	1.910	
١		 



#### MAGNEHELIC CALIBRATION CHECK

	LOW		MEI	)	HIG	H
<u>UNIT</u>	Mag	Man	Mag	Man	Mag	Man
05-24-93						
JW16 (0-1")	0.10	0.10	0.47	0.47	0.90	0.90
GF17 (0-0.5")	0.060	0.060	0.260	0.260	0.470	0.485
CA59 (0-4")	0.50	0.50	1.70	1.70	3.50	3.50
07-30-93						
R15E (0-1")	0.12	0.12	0.49	0.49	0.82	0.82
JW16 (0-1")	0.12	0.12	0.50	0.50	0.82	0.82
GF17 (0-0.5")	0.065	0.060	0.260	0.260	0.430	0.420
CA59 (0-4")	0.40	0.38	1.90	1.90	3.40	3.40
11-26-93						
R15E (0-1")	0.11	0.10	0.51	0.51	0.87	0.87
JW16 (0-1")	0.10	0.10	0.51	0.51	0.89	0.89
GF17 (0-0.5")	0.050	0.045	0.250	0.250	0.430	0.440
CA59 (0-4")	0.51	0.49	2.02	2.00	3.65	3.65

Readings in "  $\rm H_2O$ 

Reference: Oil Manometer - Dwyer #400-10 0-1" inclined, 1-10" vertical



## PRECISION INSTRUMENT REPAIR CO.

13414 WOODRUFF AVE., BELLFLOWER, CA. 90706

310 /925-6672

REPORT NO: 3916

# Certification Report

# Precision Balances and Scales

This is to certify that the balance calibrated is in compliance to US GOVERNMENT MILITARY BOOKLET, MIL-STD-45662A and that the standards used meet the compliance of NIST (NATIONAL INSTITUTE OF STANDARDS AND TECHNOLOGY)

PRODUCT ORIGINAL CONDITION: In working order.
X CALIBRATED AT IN-USE LOCATION CALIBRATED AT P.I.R. OFFICE.
TEMPERATURE: 70°F HUMIDITY: 59% STANDARD USED: Class I S/N 5943
MANUFACTURER, MODEL, SERIAL NUMBER OF PRODUCT CALIBRATED

1 TORSION TORBAL EA-1 ANALYTICAL BALANCE, S/N 156636

Calibrated to: 100mg=100.0mg

Linearity: OK Maximum Load: OK Repeteability: OK

COMPANY OR PRODUCT OWNER:

PACIFIC ENVIRONMENTAL SERVICES 13100 BROOKS, SUITE 100 BALDWIN PARK, CA. 91706 ATTN: SIYA

PO# 0640-102



TIS

Date of Calibration 12-21-93 Technichian Tom Benson License 1-0903



#### HYDROCARBON CYLINDERS VOLUME CALIBRATION 08-20-90

Cylinder #	Full, lbs	Tare, lbs	Net, 1bs	Volume, L
101	31.33	4.45	26.88	12.22
102	32.65	5.83	26.82	12.19
103	31.60	4.69	26.91	12.23
104	31.65	4.68	26.97	12.26
105	31.42	4.47	26.95	12.25
106	32.62	5.82	26.80	12.18
107	31.14	4.50	26.64	12.11
108	32.45	5.98	26.47	12.03
109	31.22	4.69	26.53	12.06
110	31.15	4.68	26.47	12.03
Supplemental	Set 07-29-92			
111	31.42	4.86	26.56	12.07
112	32.44	5.86	26.58	12.08
113	32.44	5.96	26.48	12.03
114	31.50	4.48	27.02	12.28
115	32.45	5.90	26.55	12.06
116	31.55	4.45	27.10	12.31
117	31.35	4.71	26.64	12.10
118	32.62	6.04	26.58	12.08
119	31.40	4.73	26.67	12.12
120	31.50	4.78	26.72	12.14

Note: Cylinders were filled to the rim (not including fittings) with water and weighed on a 125-lb capacity platform balance.

Water density at room temperature = 0.997 gm/cc or 0.0022 lbs/cc.



# Pitot Tube Calibration Data Sheet

Calibrated by:	na Mu	le				
Date: 12-30-93	/					
Pitot Tube I.D	5-17 A			·		
Effective Length:	5′		<u> </u>			
Pitot Tube Assembly						
Pitot Tube Openings	Damaged ?	☐ Yes	□ No			
If Yes, Explain						
$\alpha_1 = \underline{\hspace{1cm}}$	°(<10°)	$\alpha_2 = 1$	0	°(<10°)		
$\beta_1 = \underline{\qquad} \Theta$	°(<10°)	$\beta_2 = 1$	<b>D</b>	°(<10°)		
γ =•	θ =	<u> </u>	A =	0.954 •		
$z = A \sin \gamma = $	0	cm (in.)	0.32 cm	(<% in.)		
$W = A \sin \theta = $	0	cm (in.)	0.08 cm	(<1/32 in.)		
P <sub>A</sub> =	0 477	cm (in.)				
P <sub>B</sub> =	0.477	cm (in)				
D <sub>t</sub> =	0.375	cm (in.)				
Comments:						
Calibration Requ	ired ?	☐ Yes	□ No	No. 1 Table 1		

— PACIFIC ENVIRONMENTAL SERVICES, INC. —